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(54) Title: NEW USE OR CYCLOLIGNANS AND NEW CYCLOLIGNANS

(57) Abstract: The invention refers to new compounds and the use thereof as well as the use of known cyclolignans having a trans configuration of the lactone ring as inhibitors of the insulin-like growth factor-1 receptor. Said compounds can be used for treatment of IGF-1R dependent diseases, especially cancer. Especially deoxypodophyllotoxin can be used for treatment of leukemia in humans.

New use or cyclolignans and new cyclolignans

The present invention refers to new compounds as well as to the use of new and known compounds inhibiting the insulin-like growth factor-1 receptor, the IGF-1R, for treatment of IGF-1R dependent diseases, especially cancer.

BACKGROUND OF THE INVENTION

The insulin-like growth factor-1 receptor (IGF-1R) plays an important role in proliferation, protection against apoptosis and transformation of malignant cells. The IGF-1R is also important for maintaining the malignant phenotype of tumour cells, and is involved in tumour cells developing resistance to the action of anti-cancer drugs. In contrast, the IGF-1R seems not to be an absolute requirement for normal cell growth.

The IGF-1R consists of two identical extracellular alpha-subunits that are responsible for ligand binding, and two identical beta-subunits with a transmembrane domain and an intracellular tyrosine kinase domain. The ligand-receptor interaction results in phosphorylation of tyrosine residues in the tyrosine kinase domain, which spans from amino acid 973 to 1229 of the β -subunit. The major sites for phosphorylation are the clustered tyrosines at position 1131, 1135 and 1136 (LeRoith, D., et al., Endocr Rev 1995 April; 16(2), 143-63). After autophosphorylation, the receptor kinase phosphorylates intracellular proteins, like insulin receptor substrate-1 and Shc, which activate the phosphatidyl inositol-3 kinase and the mitogen-activated protein kinase signalling pathways, respectively.

Based on the pivotal role of IGF-1R in malignant cells, it becomes more and more evident that IGF-1R is a target for cancer therapy (Baserga, R., et al., Endocrine vol. 7, no. 1, 99-102, August 1997). One strategy to block IGF-1R activity is to induce

selective inhibition of the IGF-1R tyrosine kinase. However, today there are no selective inhibitors of IGF-1R available.

Drugs containing the cyclolignan podophyllotoxin has been used since centuries, and its anti-cancer properties have attracted particular interest. Undesired side effects of podophyllotoxin have, however, prevented its use as an anti-cancer drug. The mechanism for the cytotoxicity of podophyllotoxin has been attributed to its binding to beta-tubulin, leading to inhibition of microtubule assembly and mitotic arrest. The effect of podophyllotoxin on microtubules required μM concentrations in cell free systems. The *trans* configuration in the lactone ring of podophyllotoxin has been shown to be required for binding to beta-tubulin. In agreement with this, its stereoisomer picropodophyllin, which has a *cis* configuration in the lactone ring, has a 50-fold lower affinity for microtubuli and a more than 35-fold higher LD₅₀ in rats. Because of the low affinity for microtubuli of picropodophyllotoxin this compound has attracted little interest. During the last decades the major interest on podophyllotoxin derivatives has concerned etoposide, which is a ethylidene glucoside derivative of 4'-demethyl-epipodophyllotoxin. Etoposide, which has no effect on microtubules, is a DNA topoisomerase II inhibitor, and is currently being used as such in cancer therapy. A 4'-hydroxy instead of a 4'-methoxy group of such cyclolignans is an absolute requirement for them to inhibit topoisomerase II.

25

PRIOR ART

A number of synthetic tyrosine kinase inhibitors, called tyrphostins, have been studied by Párrizas, M., et al., Endocrinology 1997, Vol. 138, No. 4, 1427-1433. The IGF-1R is a member of the tyrosine kinase receptor family, which also includes the receptors of insulin, epidermal growth factor (EGF), nerve growth factor (NGF), and platelet-derived growth factor (PDGF). All of the tyrphostins active on IGF-1R cross-react with the insulin

receptor, since they are highly homologous, although two of the tyrphostins showed a moderate preference for IGF-1R. It was therefore suggested that it could be possible to design and synthesize small molecules capable of discriminating between these
5 two receptors.

Substrate competitive inhibitors of IGF-1 receptor kinase are discussed by Blum, G., et al. in Biochemistry 2000, 39, 15705-15712. A number of lead compounds for inhibitors of the isolated IGF-1R kinase are reported. The search for these compounds was
10 aided by the knowledge of the three-dimensional structure of the insulin receptor kinase domain, which is 84 % homologous to the IGF-1R kinase domain. The most potent inhibitor found was tyrphostin AG 538, with an IC₅₀ of 400 nM. However, said inhibitor also blocked the insulin receptor kinase.

15 Kanter-Lewensohn, L., et al., Molecular and Cellular Endocrinology 165 (2000), 131-137, investigated whether the cytotoxic effect of tamoxifen (TAM) on melanoma cells could depend on interference with the expression or function of the insulin-like growth factor-1 receptor. It was found that, although TAM did not
20 have a strong effect on IGF-1 binding and the expression of IGF-1R at the cell surface, at 15 microM TAM efficiently blocked tyrosine phosphorylation of the IGF-1R beta-subunit.

A connection between the IGF-1R and podophyllotoxin derivatives has never previously been made. The Chemistry of
25 Podophyllum by J.L. Hartwell et al., Fortschritte der Chemie organischer Naturstoffe 15, 1958, 83-166, gives an overview of podophyllotoxin and different derivatives thereof, commercially derived from two species of plants, *Podophyllum peltatum* and *Podophyllum emodi*. As said, the observed cytotoxic effect of
30 podophyllotoxin has been ascribed to its binding to microtubuli resulting in a mitotic block. The same effects on cells have been described for deoxypodophyllotoxin and this was suggested to be the reason why these two compounds and their corresponding 4'-demethyl

analogues could be used for treatment of psoriasis (WO 86/ 04062). However, whereas the LD₅₀ of podophyllotoxin in rats is relatively low (14 mg/kg), LD₅₀ of the deoxy derivative is strangely >15-fold higher. Other podophyllotoxin derivatives, for which LD₅₀ in rats was high, such as acetylpodophyllotoxin (185 mg/kg) and epipodophyllotoxin (>200 mg/kg) have been considered to essentially lack biological activity (Seidlova-Masinova V., et al. J Nat Cancer Inst, 18, 359-371, 1957).

Structure-activity evaluation of a number of morpholino derivatives of benzyl-benzodioxole having a structural similarity to podophyllotoxin were performed by Batra, J., et al., Biochemical Pharmacology, Vol. 35, No. 22, 4013-4018, 1986. The ability of the compounds to inhibit tubulin polymerisation was tested, but the morpholino compound most similar to podophyllotoxin was the least active in the series.

Benzyl and cinnamyl derivatives of 2,4-di-tert-butylphenol and of 1,3-benzodioxoles are known as insect chemosterilants from Jurd, L., et al., J. Agric. Food Chem. Vol. 27, No. 5, 1007-1016, 1979.

20 OBJECTS OF THE INVENTION

The object of the invention is to find new compounds and new methods for treatment of IGF-1R dependent diseases, especially cancer, by means of an inhibition of the insulin-like growth factor-1 receptor.

25

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 shows a computer model of the 12 amino acid peptide comprising the tyrosines 1131, 1135 an 1136 of the IGF-1 receptor.

Figure 2A shows the structural formulas of the compounds podophyllotoxin, deoxypodophyllotoxin, and Figure 2B shows the structural formulas of epipodophyllotoxin and acetylpodophyllo-toxin.

DESCRIPTION OF THE INVENTION

The three-dimensional structure of short peptides having the amino acid sequence of the IGF-1R tyrosine domain, including the tyrosine residues at position 1131, 1135 and 1136, were analysed 5 using a computer programme in order to find compounds having the ability to mimick the tyrosine residues and interfere with their phosphorylation. It was then discovered when using a 12-amino acid peptide that two of the three key tyrosines, that is 1135 and 1136, which have to be autophosphorylated in IGF-1R for activation, could 10 be situated as close as 0.95 nm (9.5 Å) from each other, and that the apparent angle between these groups was about 60°. The configuration of said sequence is shown in Figure 1. Such a short distance has not been observed for the corresponding tyrosines in 15 the insulin receptor. Figure 1 also depicts the space structures of podophyllotoxin and deoxypodophyllotoxin.

Molecular modelling showed that an inhibitory molecule could consist of two benzene rings separated by only one carbon atom. When a two-carbon bridge was tried, the distance between the substituents of the benzene rings was too long, about 1.3 nm (13 20 Å).

The substituents of the inhibitors corresponding to the hydroxy groups in the tyrosines were selected to be methoxy or methylenedioxy groups, since they are chemically relatively stable, i.e. they are not oxidized or phosphorylated. The distance between 25 these substituents should be about 0.95 ± 0.10 nm (9.5 ± 1.0 Å).

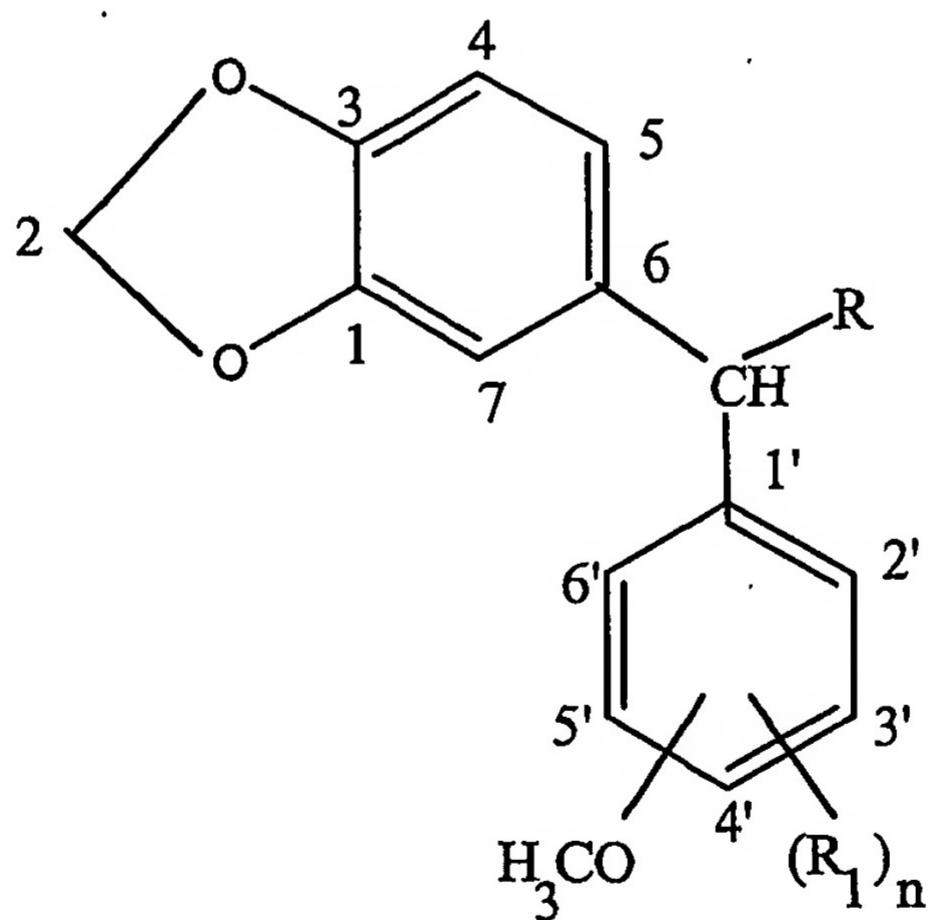
It was then surprisingly found that the two angled benzene rings of some cyclolignans, including podophyllotoxin, could mimick almost exactly the two tyrosines 1135 and 1136, indicating that 30 podophyllotoxin and derivatives could interfere with the autophosphorylation of these tyrosine residues.

In order to penetrate the receptor, an inhibitory molecule has to be small. When for instance podophyllotoxin was conjugated with a glucoside derivative, podophyllotoxin-4,6-O-benzylidene-β-D-

glucopyranoside, the effect on IGF-1R completely disappeared. Furthermore, following reduction of the lactone ring to a diol structure, the size of the molecule increased due to the reduced substituents sticking out from the molecule, resulting in a 5 dramatically reduced activity of the compounds. Increasing the size by forming methylenedioxy derivatives or acetonides of podophyllotoxindiol also resulted in compounds with little or no activity.

The inhibitor molecule also has to be relatively nonpolar, so 10 that it can freely penetrate cell membranes and the IGF-1 receptor, but sufficiently polar to be reasonably soluble in water. The polarity of the molecule is determined by the number and nature of the oxygen functions. The polarity seems to be optimal when the water solubility is between that of deoxypodophyllotoxin, i.e. 15 about 0.01 mM, and that of podophyllotoxin, about 0.1-0.2 mM. No charged or highly polar groups should be present on the molecule.

The invention refers to the use of a compound comprising the formula



wherein the distance between the carbon atom of the methylene group and the carbon atom of the methoxy group is 0.85 - 1.05 nm;

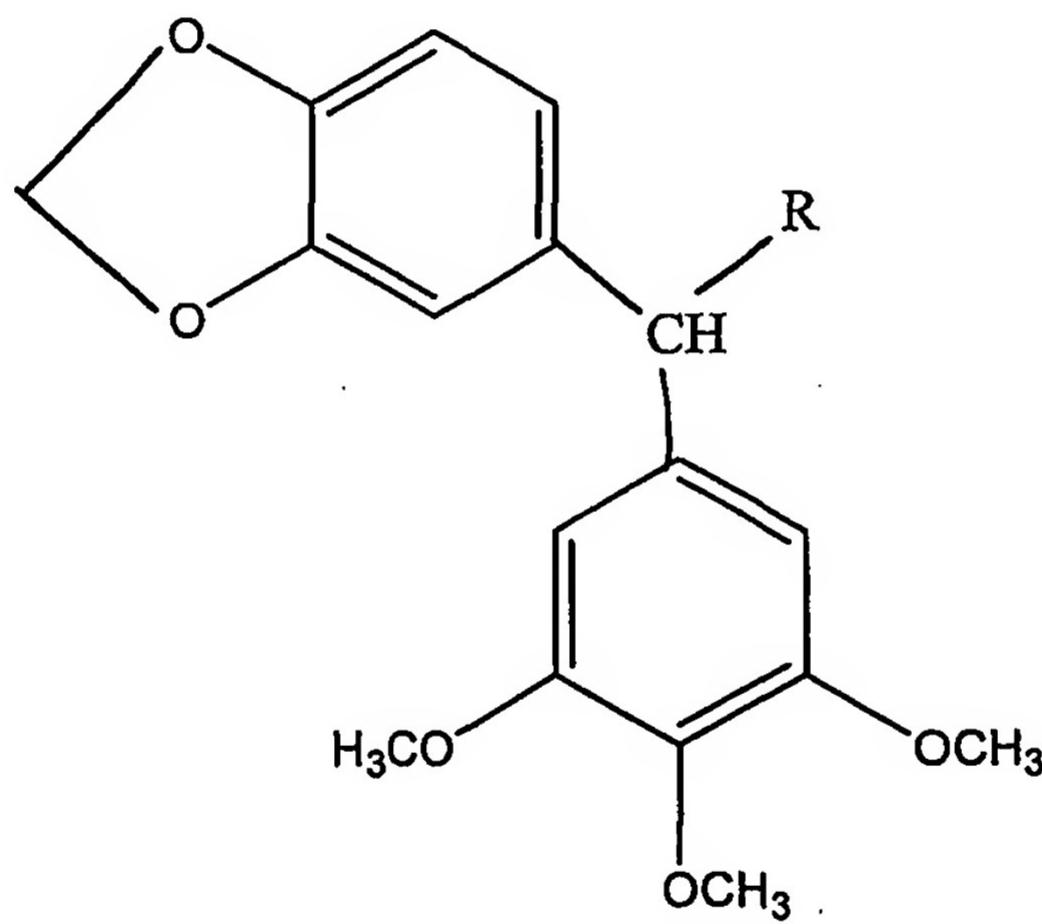
R is OH, OCH₃, OC₂H₅ or a C₁₋₅ linear or branched hydrocarbon chain, optionally having 1-3 oxygen functions, and optionally forming a bond with the carbon (number 5) in the top benzene ring.

R₁, which can be the same or different, is OH or OCH₃, and n is 0-5;

as an inhibitor of tyrosine phosphorylation of the insulin-like growth factor-1 receptor.

Oxygen functions in this context refer to hydroxy, oxo, carboxy, methoxy, methylenedioxy, lactone, ether and/or ester groups.

One group of compounds which can be used in accordance with the invention has the formula Ib

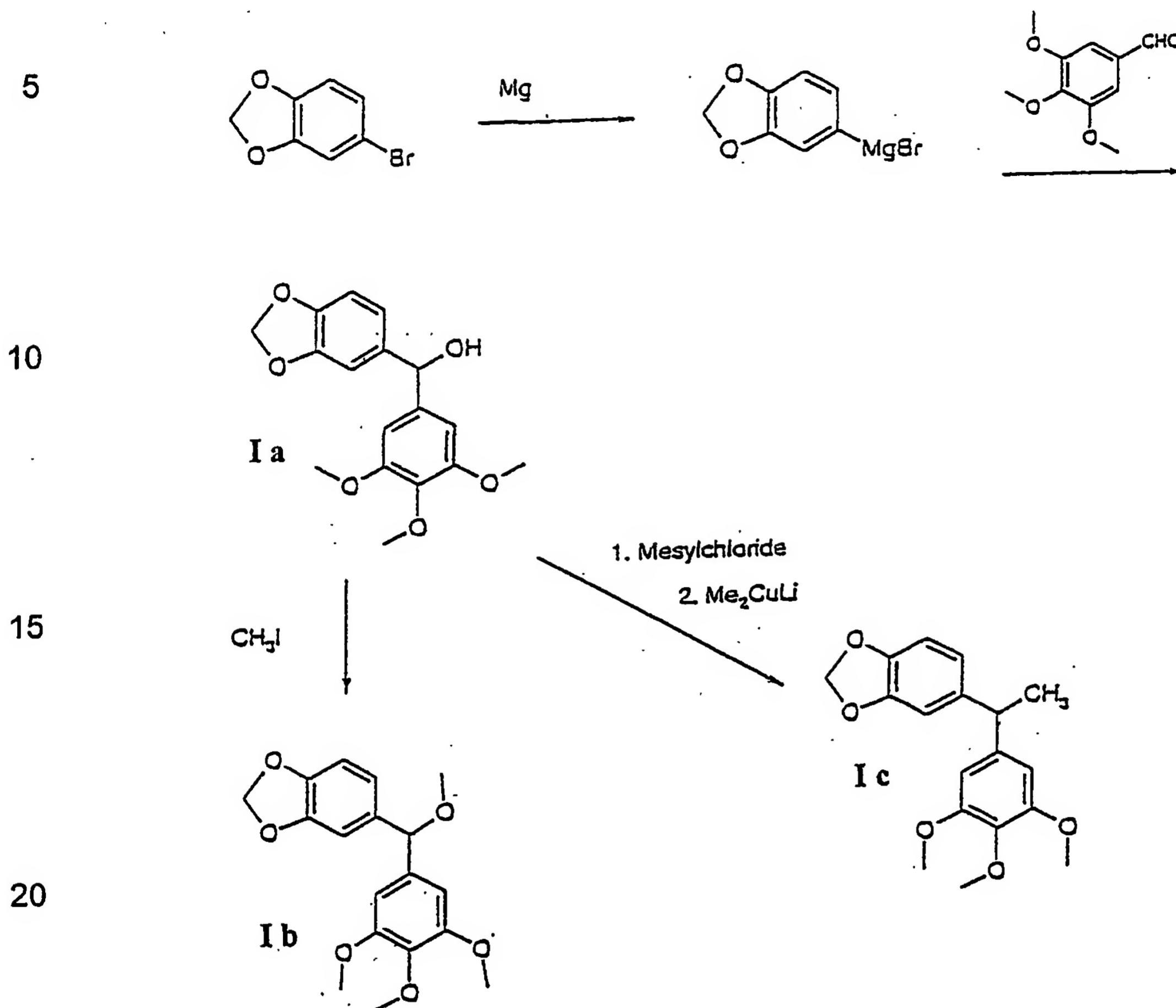


Ib

15

wherein R is OH, OCH₃, OC₂H₅, CH₃, C₂H₅, or C₂H₄OH; and R₁ and n are as defined above. The substituent R of the compounds of this group can be in the R or S position in relation to the bottom benzene ring.

Compounds of the formula I can be prepared by the following 20 representative syntheses:



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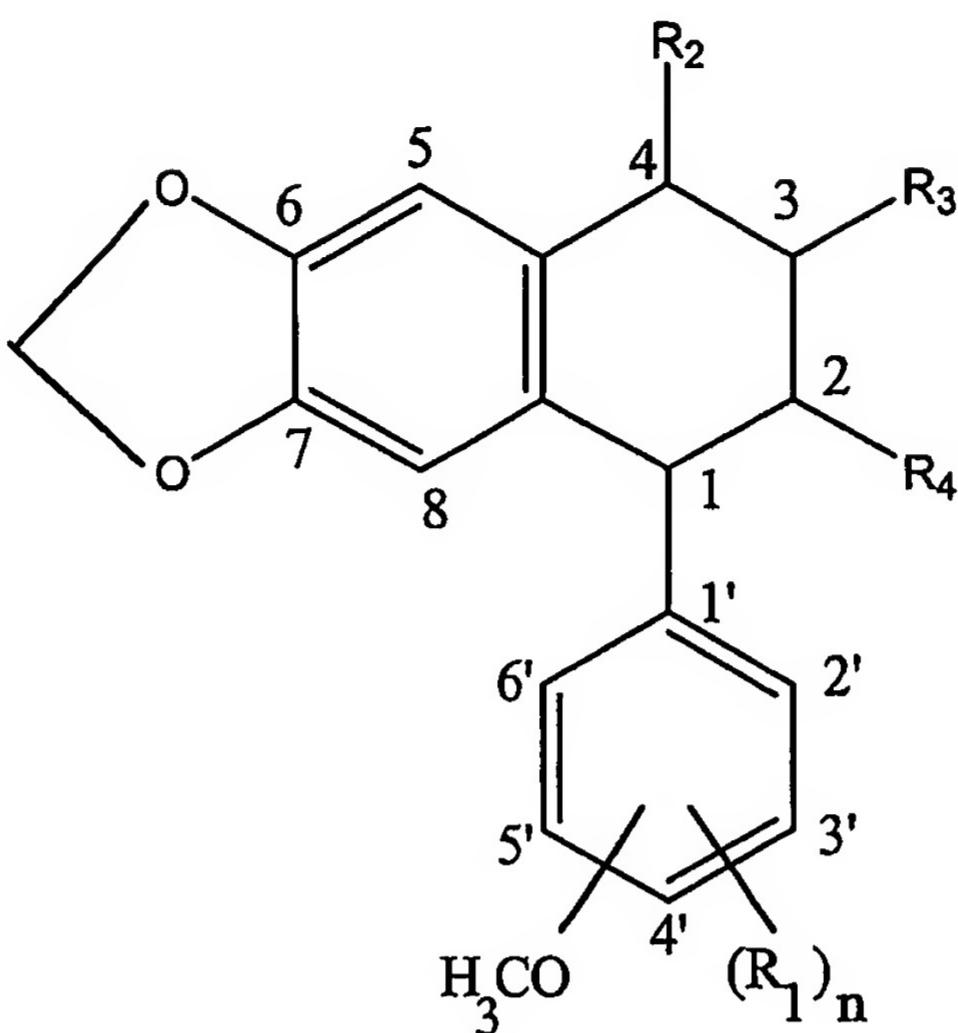
As examples of compounds which can be prepared in this way can be mentioned:

6-(3,4,5-trimethoxy-alpha-hydroxy-benzyl)-1,3-benzodioxole (Ia)

30 6-(3,4,5-trimethoxy-alpha-methoxy-benzyl)-1,3-benzodioxole (Ib)

6-(3,4,5-trimethoxy-alpha-methyl-benzyl)-1,3-benzodioxole (Ic)

Another group of compounds which can be used in accordance with the invention has the formula II



II

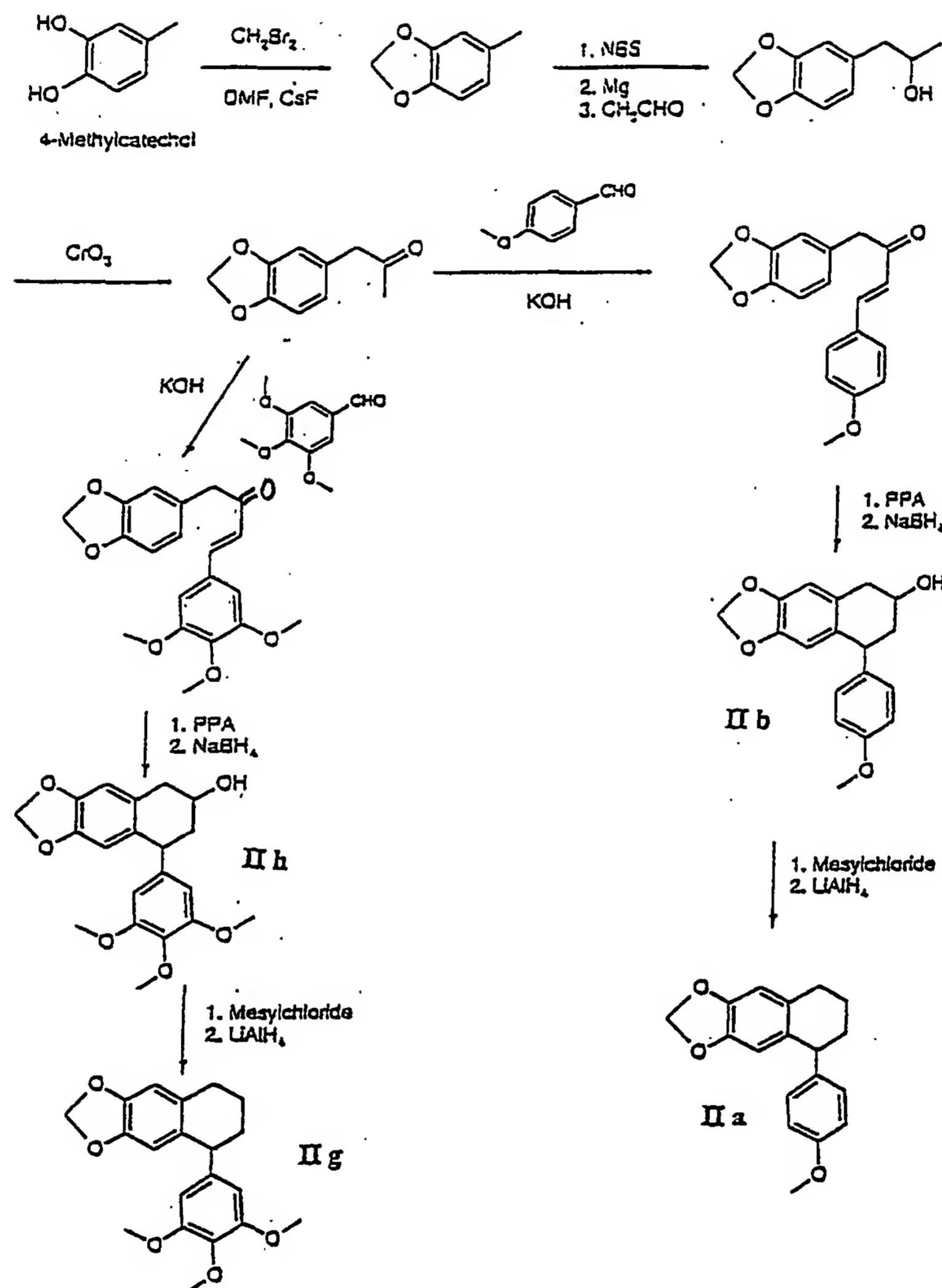
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wherein R₂, R₃ and R₄, which can be the same or different, are H, OH, O, OCH₃, OC₂H₅ or R₂ and R₃ together is a methylenedioxy group, or R₃ and R₄ together is an acetonide, carbonate or methylendioxy group; and R₁ and n are as defined above. The substituents R₂, R₃ and R₄ can, when not expressing an oxo group, be in either alpha- or beta-position. The bottom benzene ring should preferably be in the alpha-position.

Compounds of the formula II can be prepared by the following representative syntheses:

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In said schedules NBS is N-bromosuccinimide, PPA is polyphosphoric acid, TTN is thalliumnitrate trihydrate, p-TSA is p-toluene sulfonic acid, DMF is N,N-dimethylformamide, and LDA is lithium dialkylamide.

5 As examples of compounds which can be prepared in this way can be mentioned:

1-(4-methoxy-phenyl)-6,7-methylenedioxy-1,2,3,4,-
tetrahydronaphthalene (II a)

10 3-hydroxy-1-(4-methoxy-phenyl)-6,7-methylenedioxy-1,2,3,4-
tetrahydronaphthalene (II b)

1-(3,4,5-trimethoxy-phenyl)-6,7-methylenedioxy-1,2,3,4,-
tetrahydronaphthalene (II g)

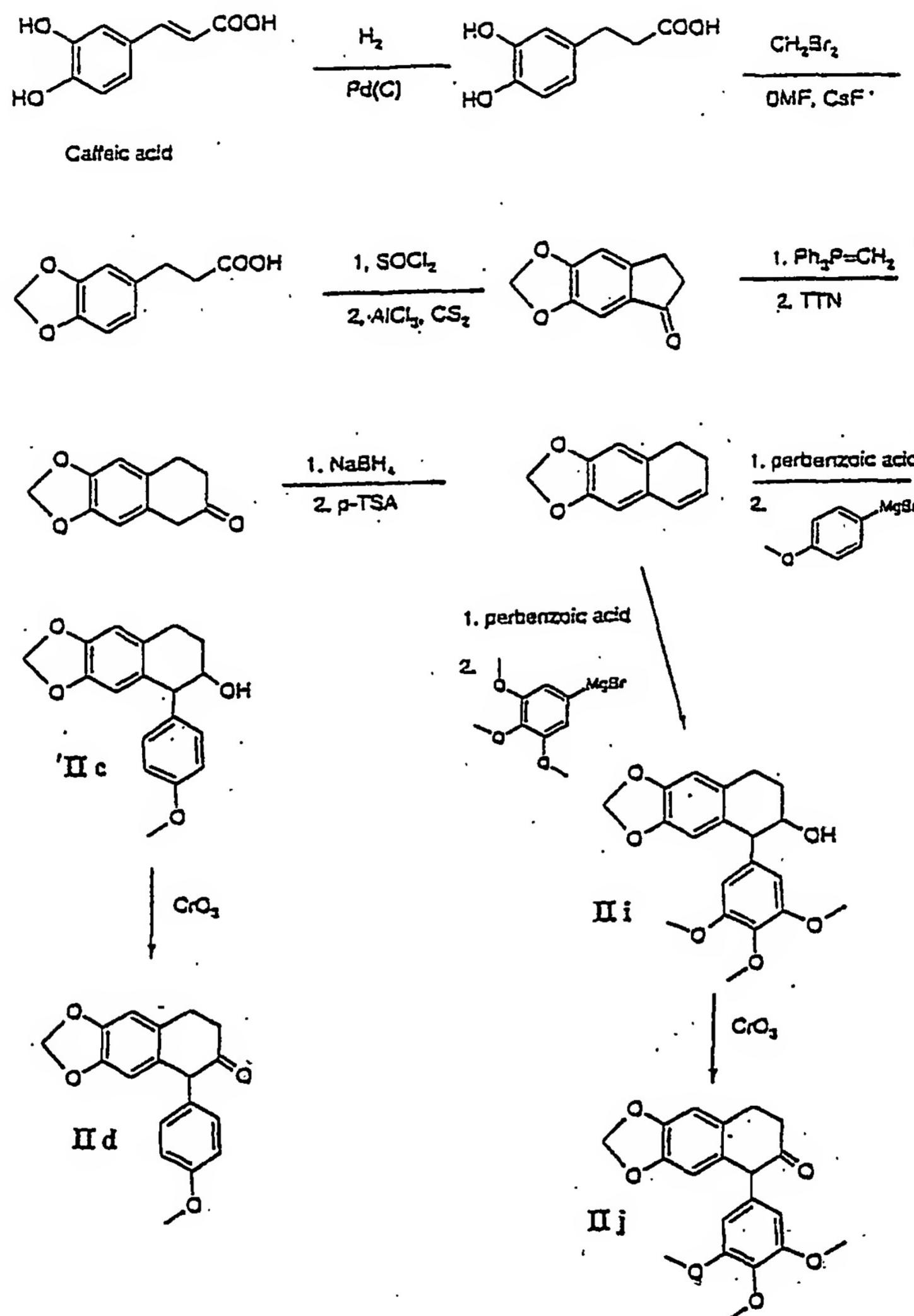
3-hydroxy-1-(3,4,5-trimethoxy-phenyl)-6,7-methylenedioxy-
1,2,3,4-tetrahydronaphthalene (II h).

15 Compounds of the formula II can also be prepared by the
following synthesis:

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As examples of compounds which can be prepared in this way can be mentioned:

2-hydroxy-1-(4-methoxy-phenyl)-6,7-methylenedioxy-1,2,3,4-

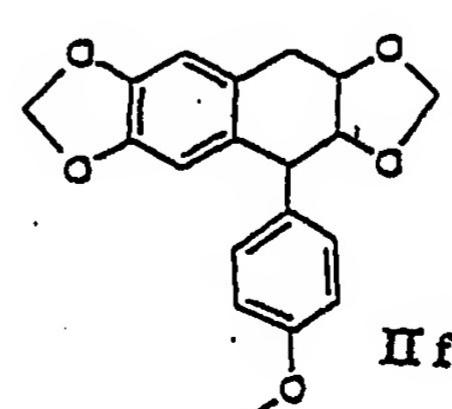
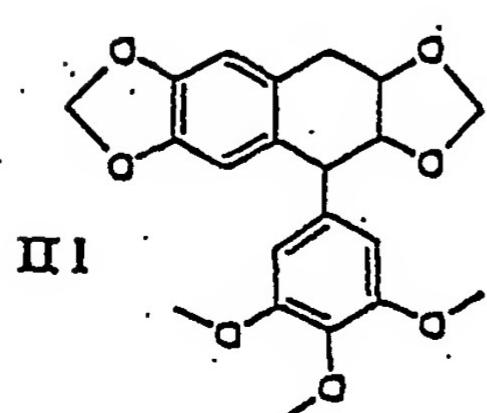
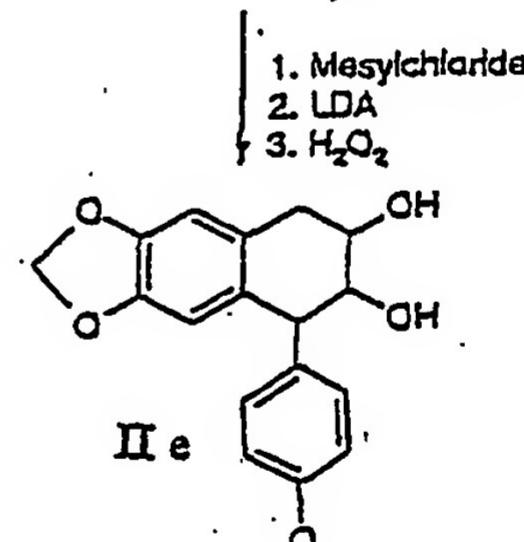
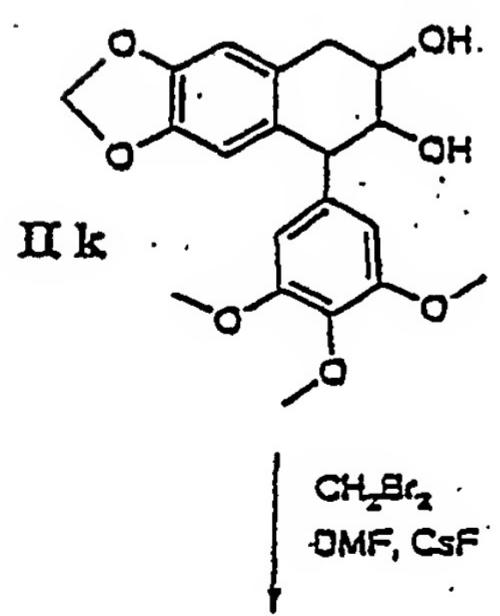
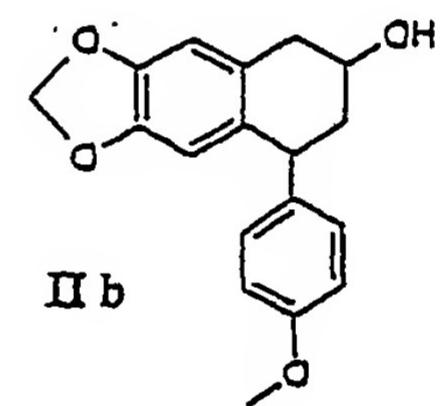
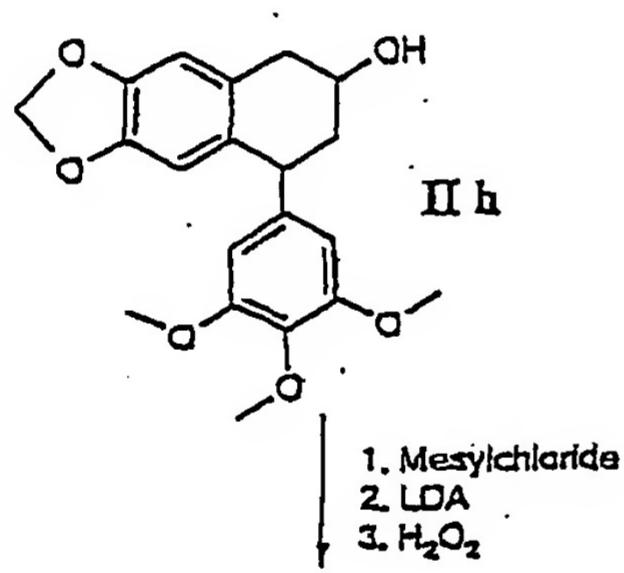
5 tetrahydronaphthalene (II c)

1-(4-methoxy-phenyl)-2-oxo-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II d)

2-hydroxy-1-(3,4,5-trimethoxy-phenyl)-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II i)

10 1-(3,4,5-trimethoxy-phenyl)-2-oxo-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II j).

Compounds of the formula II can also be prepared as follows:



1. Mesylchloride
2. LDA
3. H₂O₂

1. Mesylchloride
2. LDA
3. H₂O₂

CH₂Br₂
DMF, CsF

CH₂Br₂
DMF, CsF

As examples of compounds which can be prepared in this way can
5 be mentioned:

2,3-dihydroxy-1-(4-methoxy-phenyl)-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II e)

1-(4-methoxy-phenyl)-2,3-methylenedioxy-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II f)

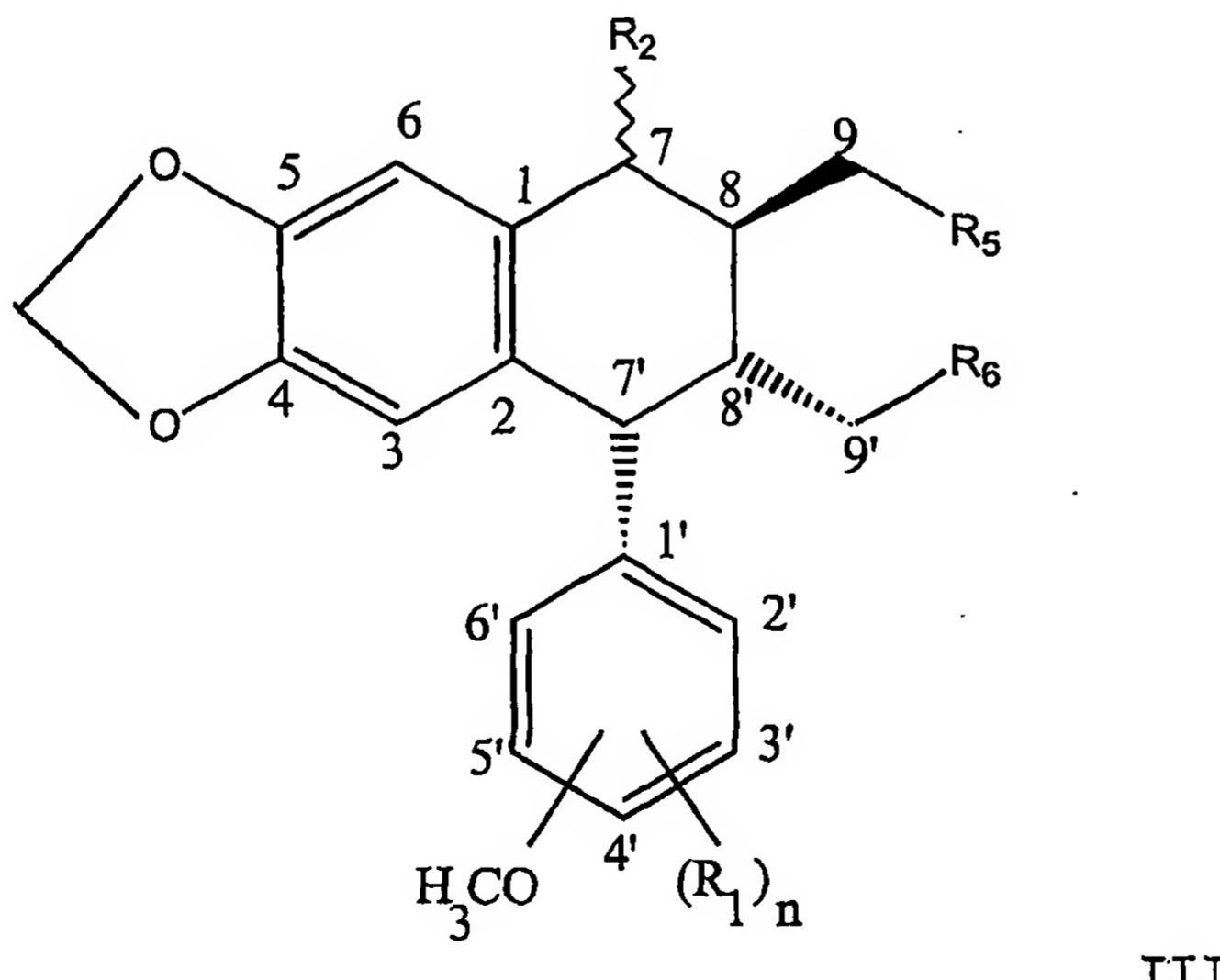
10 2,3-dihydroxy-1-(3,4,5-trimethoxy-phenyl)-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II k)

1-(3,4,5-trimethoxy-phenyl)-2,3-methylenedioxy-6,7-methylenedioxy-1,2,3,4-tetrahydronaphthalene (II l)

Details of the reaction conditions in the above syntheses
15 are described in Advanced Organic Chemistry, Jerry March (ed.), 4th edition, Wiley-Interscience Publication, New York, 1992.

Still another group of compounds which can be used in accordance with the invention are compounds of the formula III

20



wherein R₂, R₅, R₆, which can be the same or different, are H, OH, OOC₃, OOCH₂CH₃, OCH₃, or OC₂H₅, or R₅ and R₆ together is an ether or a lactone; and R₁ and n are as defined above. The substituent R₂, when being a free hydroxy group cannot be in the alpha-position, as 5 in podophyllotoxin. Other R₂ substituents, except an oxo group, can be in either alpha- or beta-position. Notably, the bottom benzene ring is in the alpha-position and there is a beta-bond between the carbons numbered 8 and 9 and an alpha-bond between the carbons numbered 8' and 9', thus they form a trans configuration, as in for 10 example deoxypodophyllotoxin and podophyllotoxin.

The invention especially refers to the use of any of the relatively non-toxic cyclolignans, such as epipodophyllotoxin, deoxypodophyllotoxin and acetylpodophyllotoxin, as an inhibitor of tyrosine autophosphorylation of the insulin growth factor-1 15 receptor, whereas the use of more cytotoxic and tissue irritating compounds, such as podophyllotoxin and 4'-demethyl-podophyllotoxin, should be avoided.

Some compounds of the formula III are naturally occurring in plants, such as deoxypodophyllotoxin and podophyllotoxin. For the 20 preparation of said substances in pure form, dried and finely ground rhizomes of e.g. *Podophyllum emodi* or *Podophyllum peltatum* are extracted with organic solvents. The extract is then filtered and concentrated on silica gel. The fractions containing the substances are collected and the latter are further purified by 25 chromatography on acid alumina and silica gel etc., and finally recrystallized. Podophyllotoxin may be used as the starting material for the syntheses of its less toxic derivatives.

Epipodophyllotoxin is readily prepared from podophyllotoxin. Five mg of the latter are dissolved in 2.5 mL of acetone. To the 30 solution is added 0.5 mL of concentrated HCl, and the mixture is boiled for 2 hours. The solution is then neutralized with aqueous NaHCO₃ (about 0.5 g in 5 mL) and following evaporation of the

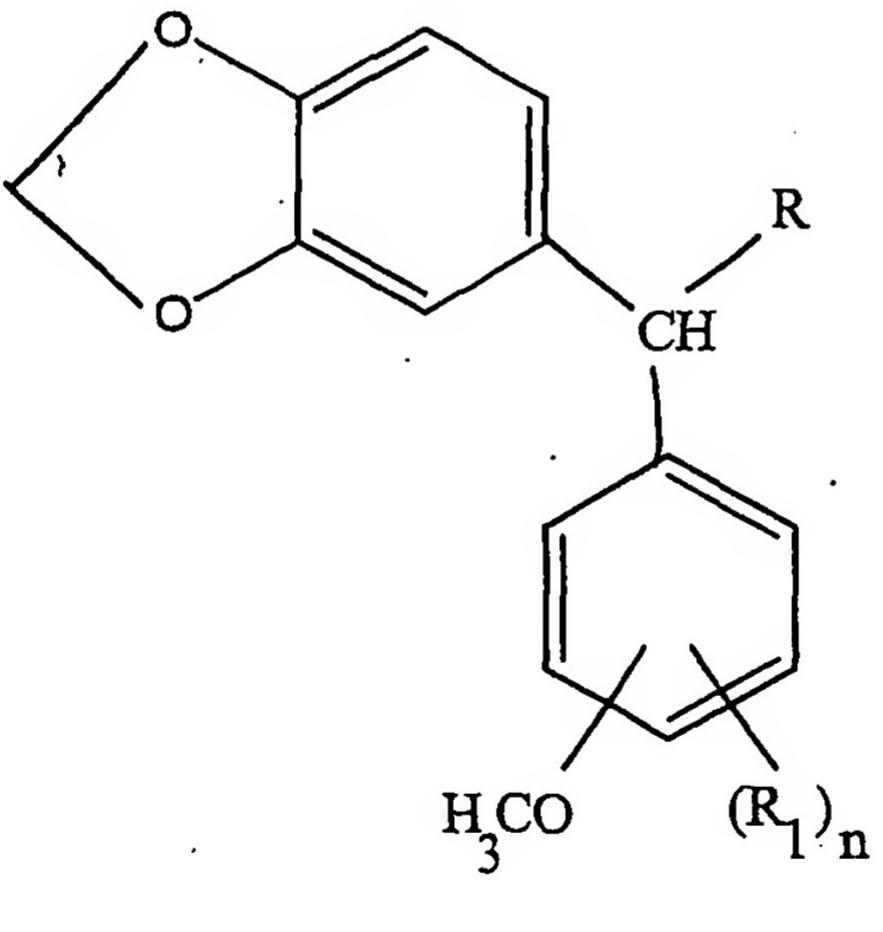
acetone, the product epipodophyllotoxin is extracted with ethyl acetate.

Acetylpodophyllotoxine (the acetate derivative of podophyllotoxin) can be prepared from podophyllotoxin by incubating 5 0.1 mg of the latter with 1 mL of acetic anhydride and 1 mL of pyridine at 50 °C for 16 hours. The reagents are then partly evaporated, 10 mL of water and 10 mL of ethyl acetate are added and the product is then extracted from the aqueous phase.

Acetonides and methylenedioxy derivatives can be prepared 10 starting from diols obtained by reducing the lactone ring of natural lignans according to standard procedures.

As additional examples of compounds of the formula III can be mentioned: podophyllotoxone, and 4'-demethyl-deoxypodophyllotoxin.

The invention also refers to the new compounds of the 15 formula I



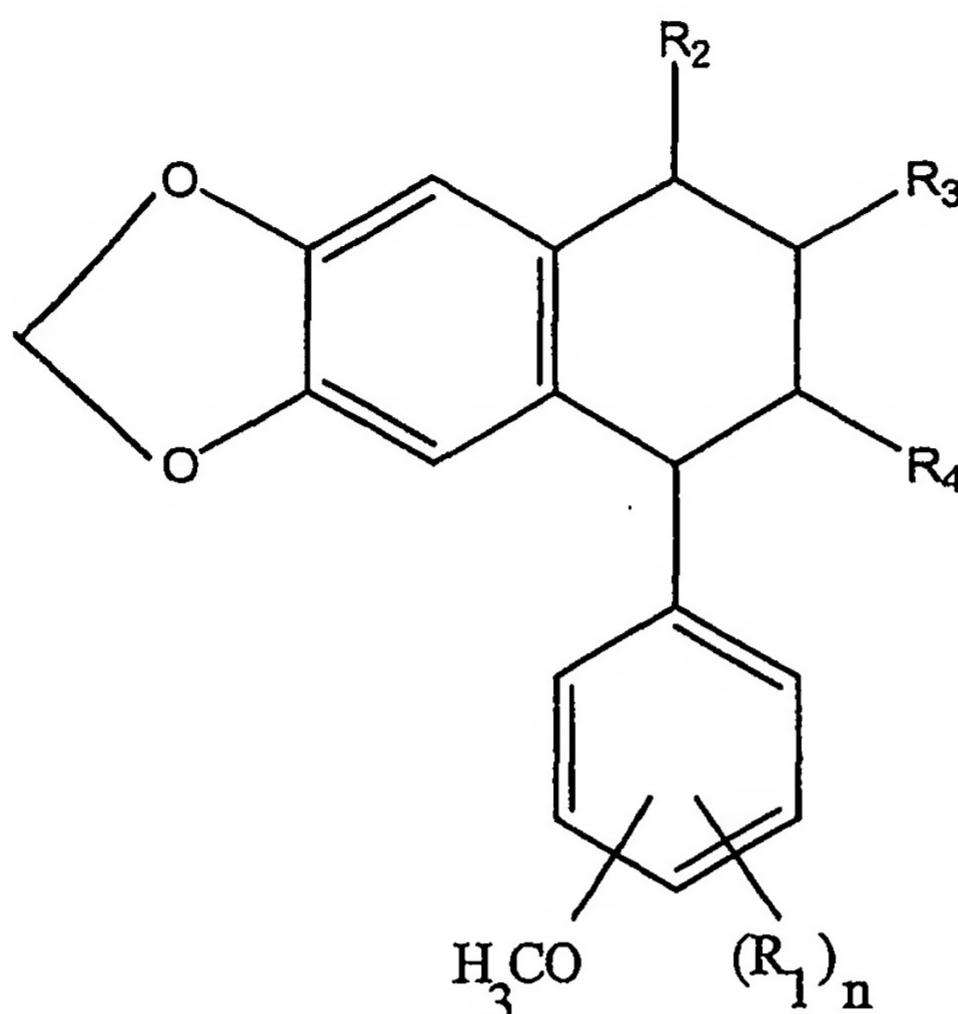
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wherein R is OH, OCH₃, OC₂H₅, CH₃, C₂H₅, or C₂H₄OH; and R₁ and n are as defined above, with the proviso that when R₁ is OCH₃ and n is 2, 20 R is not OH, that when R is a CH₃ or C₂H₅, R₁ is OH and n is 1 or 2, that when R is C₂H₅, R₁ is not OH.

The molecule of the invention should be relatively rigid, in order to keep the distance between the two substituents within the given range, that is 0.95 ± 0.10 nm (9.5 ± 1.0 Å). Forming a ring

structure of the hydrocarbon chain will prevent rotation or motion of the benzene ring, and so does lactone formation.

The invention also refers to the new compounds of the formula II



5

II

wherein R₂, R₃ and R₄, which can be the same or different, are H, OH, O, OCH₃, or R₂ and R₃ together is a methylenedioxy group, or R₃ and R₄ together is an acetonide, carbonate or methylendioxy group; and R₁ and n are as defined above.

10 To design an inhibitor of the IGF-1R tyrosine kinase for therapeutic purposes it is of critical importance that the inhibitor does not cross-react with the insulin receptor kinase, which is highly homologous to the IGF-1R. Co-inhibition of the insulin receptor will lead to a diabetogenic response in-vivo. This
 15 response comprises a very serious side effect, which cannot be overcome by insulin treatment since the receptor kinase is being blocked. We have demonstrated that podophyllotoxin and derivatives, which are much more potent IGF-1R inhibitors than the tyrophostin-based compounds, do not interfere with the insulin receptor
 20 tyrosine kinase at all. Neither do they interfere with tyrosine

phosphorylation of the receptors of epidermal growth factor, platelet-derived growth factor or fibroblast growth factor.

Podophyllotoxin has for long been implicated in cancer therapy, but in the way it was administered to patients it produced unacceptable side effects. The anti-cancer effect, as well as the side effects, was attributed to inhibition of microtubule assembly and mitotic block. It has now been demonstrated that podophyllotoxin and some of its less toxic analogues are very potent and specific inhibitors of tyrosine phosphorylation of the insulin-like growth factor-1 receptor, which plays a pivotal role as a survival factor in cancer cells. Compared to the anti-microtubule effect of podophyllotoxin, a 100-fold lower concentrations were sufficient to inactivate the IGF-1R. Of utmost importance is that podophyllotoxin and analogues do not inhibit the insulin receptor, which is highly homologous to IGF-1R. Moreover, they do not inhibit other major growth factor receptor kinases either.

Relatively nontoxic compounds of the formula I can be used for treatment of IGF-1R dependent diseases, such as cancer, arteriosclerosis, including prevention of restenosis of the coronary arteries after vascular surgery, psoriasis and acromegaly.

A pharmaceutical composition comprising a compound of the formula I in combination with a physiologically acceptable carrier and optional additives can be administered to a patient by any suitable route, such as parenterally, preferably by intravenous infusion, or topically, for instance by a patch.

The invention refers to the new compounds of the formula I or II for use as a medicament, and especially for the preparation of a medicament for treatment of cancer.

The results of the biological experiments suggest that submicromolar concentrations of podophyllotoxin, or of less toxic analogues such as deoxypodophyllotoxin or epipodophyllotoxin, can be sufficient to cause tumour cell death. However, it is believed

that it is important to keep a constant plasma concentration of the inhibitors over lengthy periods, to allow them to continuously saturate all IGF-1Rs, and in this way eventually kill as many malignant cells as possible. Therefore, continuous infusion of 5 podophyllotoxin derivatives, in connection with monitoring the plasma concentration, may be the strategy of treatment instead of repetitive (e.g. daily) injections, which may lead to repeated reactivations of IGF-1R between the treatments.

The invention consequently also refers to a method of 10 treatment of a cancer in a mammal, comprising the steps of administrating a pharmaceutical composition, containing a compound having the formula I in combination with a physiologically acceptable carrier, by constant infusion to a patient suffering from a tumour, controlling the plasma level of the compound, and 15 adjusting the rate of infusion to keep the plasma level between 0.05 and 5.0 μM (depending on the general toxicity of the compound), for a period of time being sufficient for the tumour to be retarded or to disappear.

In case of tumours not completely dependent on IGF-1R, the 20 compounds of the invention can be useful to sensitise the tumour cells to the effects of other anti-cancer drugs.

EXPERIMENTAL

Materials

25 Chemicals

Cell culture reagents, that is media, fetal calf serum and antibiotics, were purchased from Gibco, Sweden. All other chemicals unless stated otherwise were from Sigma (St. Louis. MO, USA). A mouse monoclonal antibody against phosphotyrosine (PY99) and a 30 polyclonal antibody against α -subunit of IGF-1R (N20) were obtained from Santa Cruz Biotechnology Inc (Santa Cruz, CA, USA). A monoclonal antibody against the α -subunit of IGF-1R (IR-3) was purchased from Oncogene Science (N.Y., USA). Deoxypodophyllotoxin

and podophyllotoxin (99.97% purity), and acetylpodophyllotoxin, podophyllotoxone and 4'-demethylpodophyllotoxin (>95 % purity) were obtained as gifts from Analytecon SA, Switzerland.

Cell cultures

5 The human malignant melanoma cell lines SK-MEL-2, SK-MEL-5 and SK-MEL-28, the prostatic carcinoma cell line PC-3, and the breast cancer cell line MCF-7 were from the American Tissue Culture Collection, USA. The malignant melanoma cell lines BE, and FM55 were obtained from Professor R Kiessling, CCK, Karolinska Hospital, 10 Stockholm, Sweden. The R- and P6 cell lines were gifts from Professor R. Baserga, Thomas Jefferson University, Philadelphia, PA, USA. All cell lines were cultured in Minimal Essential Medium containing 10% faetal bovine serum, glutamine, 1% benzylpenicillin and streptomycin. The cells were grown in monolayers in tissue 15 culture flasks maintained at 95 % air/5 % CO₂ atmosphere at 37°C in a humidified incubator. For the experiments cells were cultured in either 35-mm or 60-mm plastic dishes or 96-well plastic plates. The experiments were initiated under subconfluent growth conditions.

 The human chronic myeloid leukemia K562/S and K562/Vcr30 lines 20 and the acute myeloid leukemia cell lines HL60/0 and HL60/Nov were obtained from ATCC. The K562/S and HL60/0 are wild type (non-resistant) cells, whereas K562/Vcr30 and HL60/Nov are cytostatic-resistant sublines. All leukemia cell lines were cultured in RPMI 1640 medium supplemented with 10% fetal bovine serum and with 2 mM 25 glutamine, 1 % benzyl-penicillin and streptomycin. The cells were grown in tissue culture flasks maintained at 95 % air/5% CO₂ atmosphere at 37°C in a humidified incubator. For the experiments 25,000 cells were cultured in 60-mm plastic dishes or 96-well plastic plates. The experiments on leukemia cells were performed in 30 collaboration with Associate professor Sigurd Vitols, Department of Pharmacology, Karolinska Hospital (Stockholm, Sweden).

Methods

Assay of cell growth and survival

Cell proliferation kit II (Roche Inc.) is based on colorimetric change of the yellow tetrazolium salt XTT in orange formazan dye by the respiratory chain of viable cells (Roehm, NW, et al., J Immunol Methods 142:257-265, 1991). Cells seeded at a 5 concentration of 5000/well in 100 µl medium in a 96-well plate were treated with different drugs in the given concentration. After 24 or 48 h the cells were incubated, according to the manufacturer's protocol, with XTT labelling mixture. After 4 h the formazan dye is quantified using a scanning multiwell spectrophotometer with a 495-10 nm filter. The absorbance is directly correlated with number of viable cells. The standard absorbance curve was drawn by means of untreated cells seeded at a concentration of from 1000 to 10 000 cells/ well with an increasing rate of 1000 cells/ well. All standards and experiments were performed in triplicates.

15 Immunoprecipitation and determination of protein content

The isolated cells were lysed in 10 ml ice-cold PBSTDs containing protease inhibitors (Carlberg, M., et al., J Biol Chem 271:17453-17462, 1996). 50 µl protein A or G agarose was added in 1 ml sample and incubated for 15 min at 4°C on an orbital shaker. 20 After centrifugation for 10 min at 10,000 r/min at 4°C the supernatant was saved. The protein content was determined by a dye-binding assay with a reagent purchased from Bio-Rad. Bovine serum albumin was used as a standard. 15 µl Protein G Plus agarose and 5 µl anti-IGF-1R were added. After a 3 h incubation at 4°C on an 25 orbital shaker the precipitate was collected by pulse centrifugation in a micro centrifuge at 14,000xg for 10 s. The supernatant was discarded and the pellet was washed 3 times with PBSTDs.

Sodium dodecyl sulphate polyacrylamide gel electrophoresis

30 (SDS-PAGE)

Protein samples were solved in a 2x-sample buffer containing Laemmli buffer and 0.5% methanol and boiled for 5 min at 96°C. Samples were separated by SDS-PAGE with a 4% stacking gel and 7.5%

separation gel. Molecular weight markers (Bio Rad, Sweden) were run simultaneously in all experiments.

Western blotting

Following SDS-PAGE the proteins were transferred overnight to 5 nitro-cellulose membranes (Hybond, Amersham, UK) and then blocked for 1 h at room temperature in a solution of 4% skimmed milk powder and 0.02% Tween 20 in PBS, pH 7.5. Incubations with the primary antibodies were performed for 1 h at room temperature, followed by 3 washes with PBS with Tween and incubation with the second 10 antibody for 1 h room temperature. After another 3 washes the membranes were incubated with Streptavidin-labelled horseradish peroxidase for 30 min and then detected using Amersham ECL system (Amersham, UK). The films were scanned by Fluor-S (BioRad).

Assay of IGF-1R autophosphorylation in vitro

15 IGF-1R tyrosine autophosphorylation was analysed by a sandwich ELISA assay. Briefly, 96-well plates (Immunolon, Nunc) were coated overnight at 4 °C with 1 µg/well of the monoclonal antibody Ab-5 (LabVision) to the IGF-1R beta subunit. The plates were blocked with 1% BSA in PBS Tween for 1 h, then 80 g/well of total protein 20 lysate from the P6 cell line was added. As a negative control was used total protein lysate from R-cell line. The investigated compounds were added in tyrosine kinase buffer without ATP at room temperature for 30 min, prior to kinase activation with ATP. Kinase assay was performed using the Sigma kit. After spectrophotometry 25 the IC₅₀ values of inhibitors were determined using the Regression function of Statistica program.

Experiment 1. Effect of podophyllotoxin and other biologically active phenolic compounds on phosphorylation of IGF-1R in cultured 30 melanoma cells

Melanoma cells (line FM55) were seeded in 6-cm dishes, at a concentration of 10,000 cells/cm² in Minimal Essential Medium supplemented with 10% fetal calf serum (FCS). When the cells

reached a concentration of 65,000 cells/cm², they were treated with genistein, tamoxifen, quercetin and podophyllotoxin to a final concentration of 0, 1, 15 or 60 µM in the culture medium for 1 h. Treatment with 0 µM represents untreated controls. The cells were 5 then isolated and subjected to immuno-precipitation of the IGF-1R. The immunoprecipitates, containing purified IGF-1R, were fractionated by gel electrophoresis. Phosphorylation of IGF-1R was detected by an anti-phosphotyrosine antibody using Western blotting. The obtained signals represent phosphorylated IGF-1R and 10 the intensity of signals represents amounts of phosphorylated IGF-1R. Details of the methods used are described above. The intensities are quantified by a scanner, which measures the optical density (OD) of the signals. For the control cells the OD is set at 100%. The blank (OD 0%) represents the background. The results 15 given in Table 1 below are mean values of 3 experiments.

Table 1. Level of IGF-1R phosphorylation in intact cells

Compound	1 µM	15 µM	60 µM
Genistein	100	96	35
Tamoxifen	95	20	10
Quercetin	100	105	96
Podophyllotoxin	8	4	2

The results show that podophyllotoxin almost completely blocks 20 IGF-1R phosphorylation at all three concentrations., whereas genistein only has a partial inhibitory effect at 60 µM and quercetin has no effect at all.

Experiment 2. Effect of podophyllotoxin derivatives on
autophosphorylation of IGF-1R in cultured melanoma cells

FM 55 melanoma cells were cultured in the same way as described in Experiment 1. When reaching a density of 65,000 cells/cm² in the dishes, they were treated for 1 h with 0.05 µM podophyllotoxin,

deoxypodophyllotoxin, acetylpodophyllotoxin, epipodophyllotoxin, 4'-demethyl-podophyllotoxin and podophyllotoxone. The cells were then harvested for assay and quantification of IGF-1R autophosphorylation as described above. The values shown in Table 2
5 represent means of 3 experiments.

Table 2. Inhibitory effect on IGF-1R autophosphoryl-ation in intact cells in relation to podophyllotoxin

Compound	Relative potency
Podophyllotoxin	1
Deoxypodophyllotoxin	0.8
Acetylpodophyllotoxin	1.3
Epipodophyllotoxin	0.5
4'-demethylpodophyllotoxin	0.5
Podophyllotoxone	0.3

10 The results show that acetylpodophyllotoxin, podophyllotoxin, and deoxypodophyllotoxin are potent inhibitors of IGF-1R phosphorylation.
15

Experiment 3. Dose-response effects of podophyllotoxin and deoxypodophyllotoxin on viability of solid tumour cells

5 different types of cell lines were seeded in 96-well plates (medium volume in a well was 100 µl), at a concentration of 10,000 cells/cm² in Minimal Essential Medium supplemented with fetal calf serum. When the cells had reached a concentration of 65,000
20 cells/cm², they were treated with different doses of podophyllotoxin and deoxypodophyllotoxin for 48 h. Cell viability was then assayed (see above). IC50 values for each inhibitor and cell line, calculated as the concentration, resulting in a 50 % decrease in cell survival, are shown below. The results are based
25 on 4 different experiments.

Table 3. IC50 (μ M) for cell viability

Cell line	Origin	Podophyllotoxin	Deoxypodophyllotoxin
SK-MEL-28	melanoma	0.05	0.04
BE	melanoma	0.05	nd
FM55	melanoma	0.04	0.04
MCF-7	Breast cancer	0.07	0.03
PC-3	prostate cancer	0.06	Nd

• nd, not determined

This shows that podophyllotoxin and deoxypodophyllotoxin are both
5 very potent inhibitors of tumor cell viability.

Experiment 4. Dose-response effects of different podophyllotoxin analogues

FM55 melanoma cells were cultured in the same way as described
10 in Experiment 3 and were treated with different doses of
podophyllotoxin analogues as described in Experiment 3. The results
(IC50 values) are given in the following Table 4.

Table 4. IC50 (μ M) for viability of FM55 cells

Compound	IC50
Podophyllotoxin	0.05
Deoxypodophyllotoxin	0.04
Acetylpodophyllotoxin	0.03
4'-demethylpodophyllotoxin	0.04

15

The result shows that the tested analogues were all potent.

Experiment 5. Dose-response effects of podophyllotoxin analogues on viability of leukemia cell

20 The leukemia cell lines K562/S, K562/Vcr 30, HL60/0 and HL60/Nov were proven to express the IGF-1R. This was assayed by Western blotting analysis, as described in Methods and Experiment 1

and 2. The 4 leukemia cell lines were seeded in 96-well plates (medium volume in a well was 100 µl), in RPMI40 medium supplemented with fetal calf serum. After 24 h podophyllotoxin, deoxypodophyllotoxin and other derivatives were added at different 5 concentrations for 72 h. Cell viability was then assayed (see above). IC₅₀ values for each inhibitor and cell lines are shown below (Table 5). The results are based on 3 different experiments.

Table 5. IC₅₀ (nM) for viability of leukemia cell lines

	K562/S	K562/Vcr30	HL60	HL60/Nov
Podophyllotoxin	4	7	3	2
Deoxypodophyllotoxin	3	4	3	2
Acetylpodophyllotoxin	80	140	48	46
Epipodophyllotoxin	195	450	127	90
Podophyllotoxone	>500	>500	>500	>500
4'-Demethyl-podophyllotoxin	22	50	20	18

10

The results demonstrate that deoxypodophyllotoxin has an unexpected and exceptionally strong cytotoxic effect on human leukemia cells. Such an effect can not be explained solely by its action on IGF-1R. The effects of acetylpodophyllotoxin and epipodophyllotoxin are as 15 expected for an IGF-1R inhibitor.

CONCLUSION

It has been demonstrated that certain cyclolignans such as podophyllotoxin and some analogues deoxypodophyllotoxin are highly 20 specific and potent inhibitors of the IGF-1R tyrosine kinase, as assayed in intact cells. When administered to intact cells the EC₅₀ value was as low as 0.02-0.06 µM.

Podophyllotoxin-induced inactivation of the insulin-like growth factor-1 receptor caused extensive cell death in malignant 25 cells, whereas cells devoid of insulin-like growth factor-1 receptors were resistant. The non-toxic derivative picropodophyllin

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was equipotent to podophyllotoxin in inhibiting the insulin-like growth factor-1 receptor activity and inducing cell death. This new mechanism of podophyllotoxin and derivatives may be useful in therapy of cancer and other IGF-1R dependent diseases.

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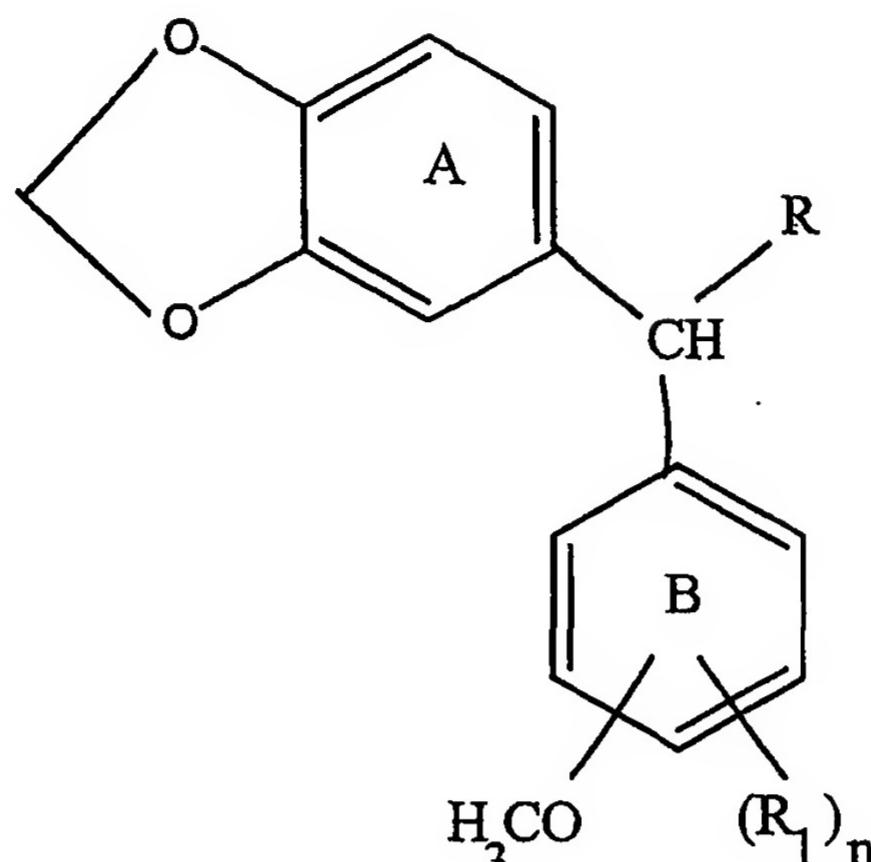
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CLAIMS

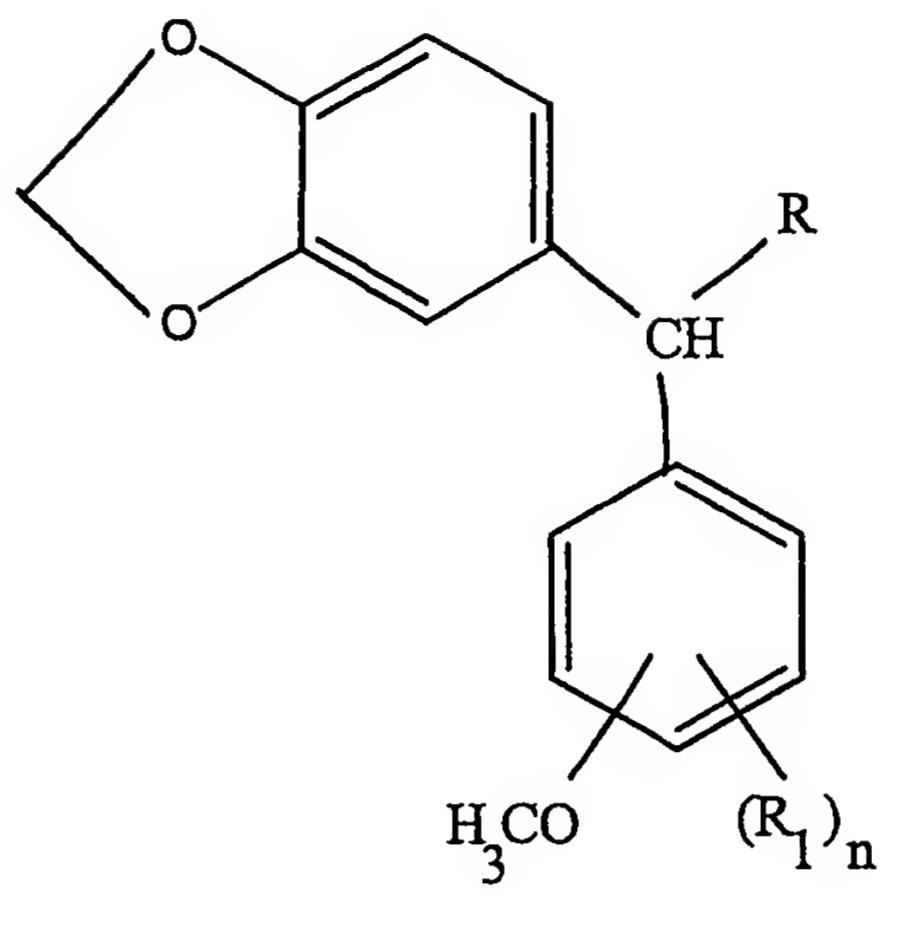
1. Use of a compound having the formula I



5

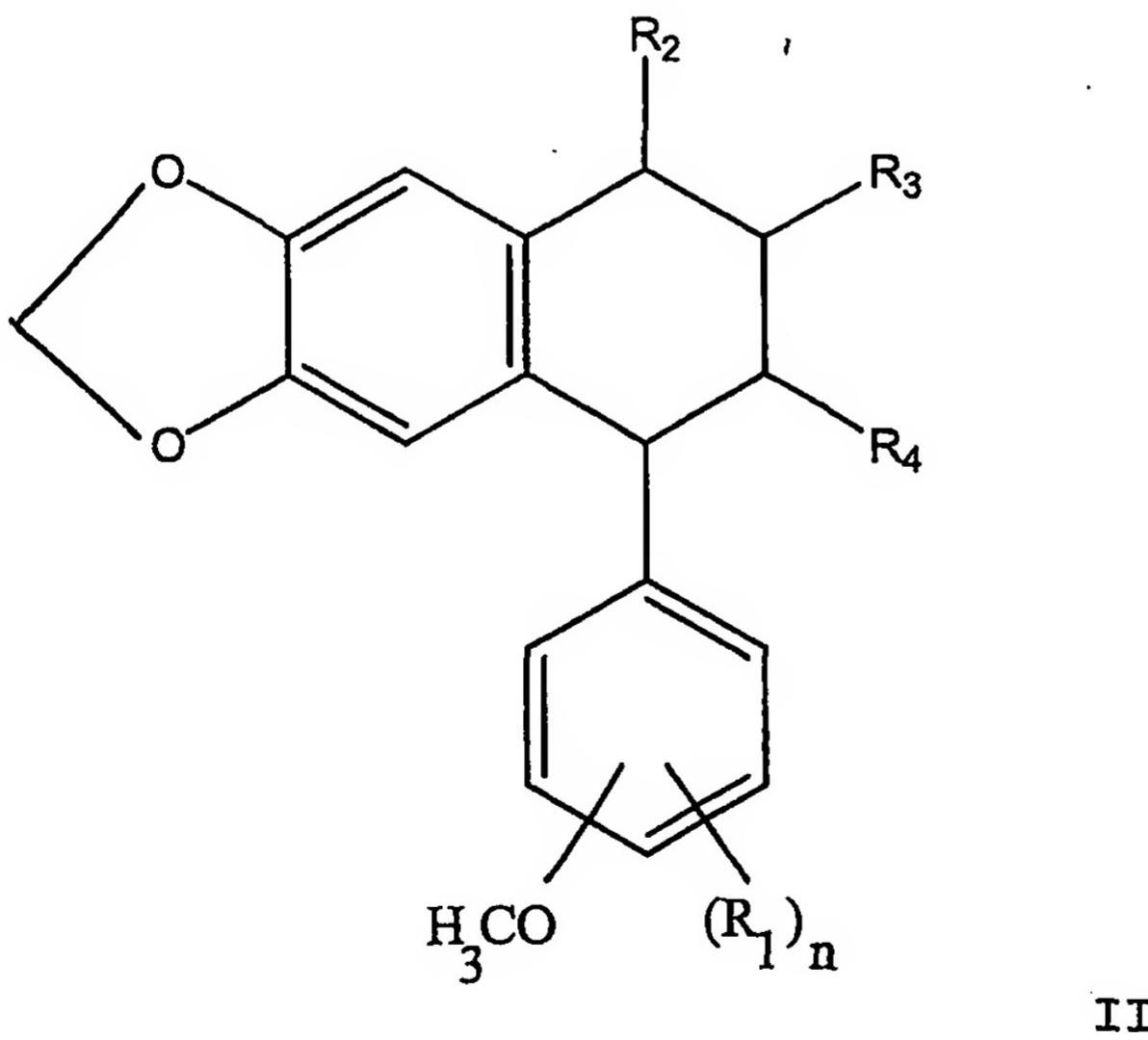
wherein the distance between the carbon atom of the methylene group and the carbon atom of the methoxy group is 0.85-1.05 nm;
 R is OH, OCH₃, OC₂H₅ or a C₁₋₅ linear or branched hydrocarbon chain
 10 with optionally a double bond, optionally having 1-3 oxygen functions, and optionally forming a bond with a carbon atom in the benzene ring A;
 R₁, which can be the same or different, is OH or OCH₃, and n is 0-2;
 15 as an inhibitor of tyrosine phosphorylation of the insulin-like growth factor-1 receptor.

2. Use according to claim 1 of a compound having the formula I



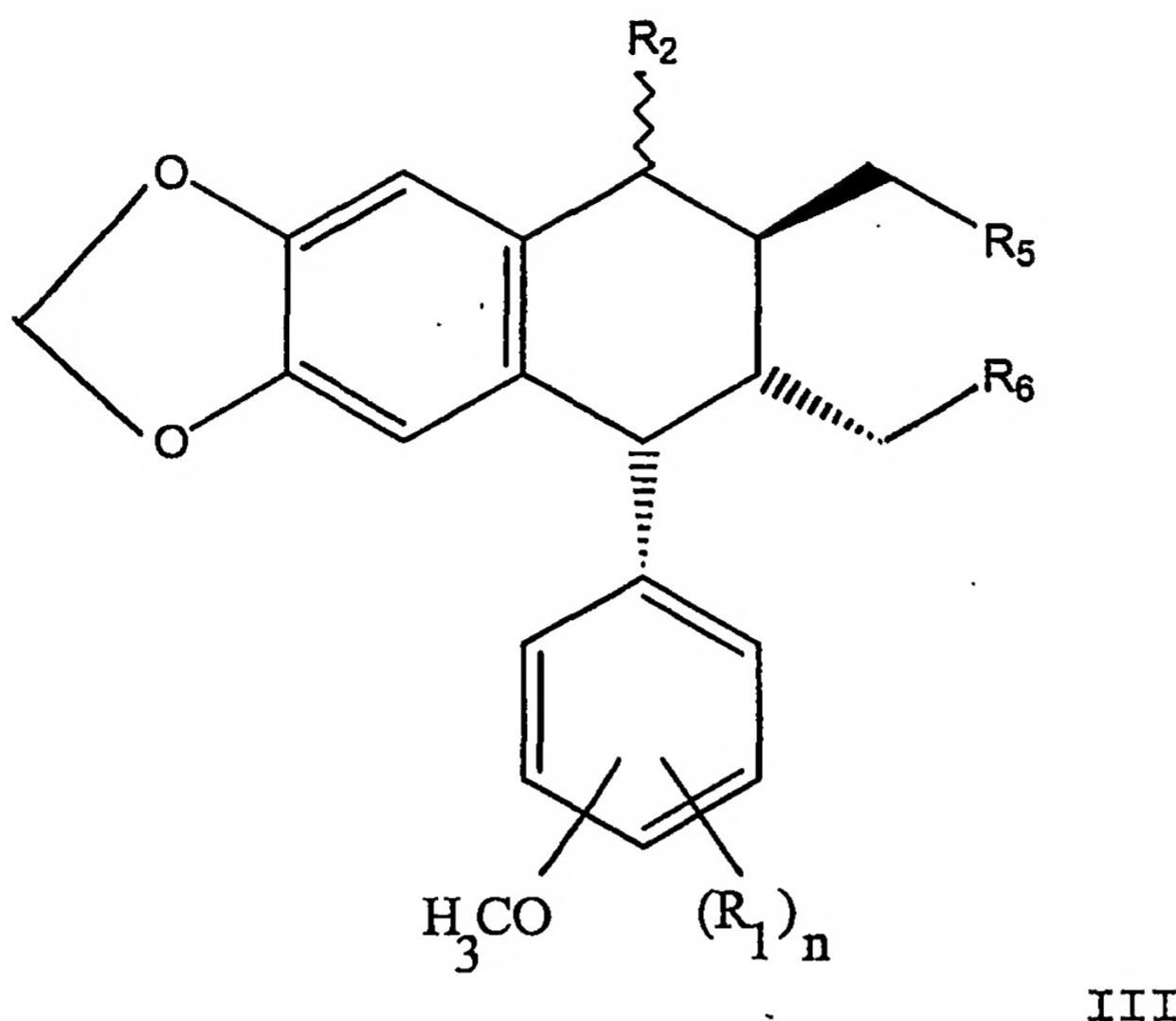
wherein R is OH, OCH₃, OC₂H₅, CH₃, C₂H₅, or C₂H₄OH; and R₁ and n are
5 as defined in claim 1.

3. Use according to claim 1 of a compound of the formula II



wherein R₂, R₃ and R₄, which can be the same or different, are H,
10 OH, O, OCH₃, or R₂ and R₃ together is a methylenedioxy group, or R₃ and R₄ together is an acetonide, carbonate or methylendioxy group; and R₁ and n are as defined in claim 1.

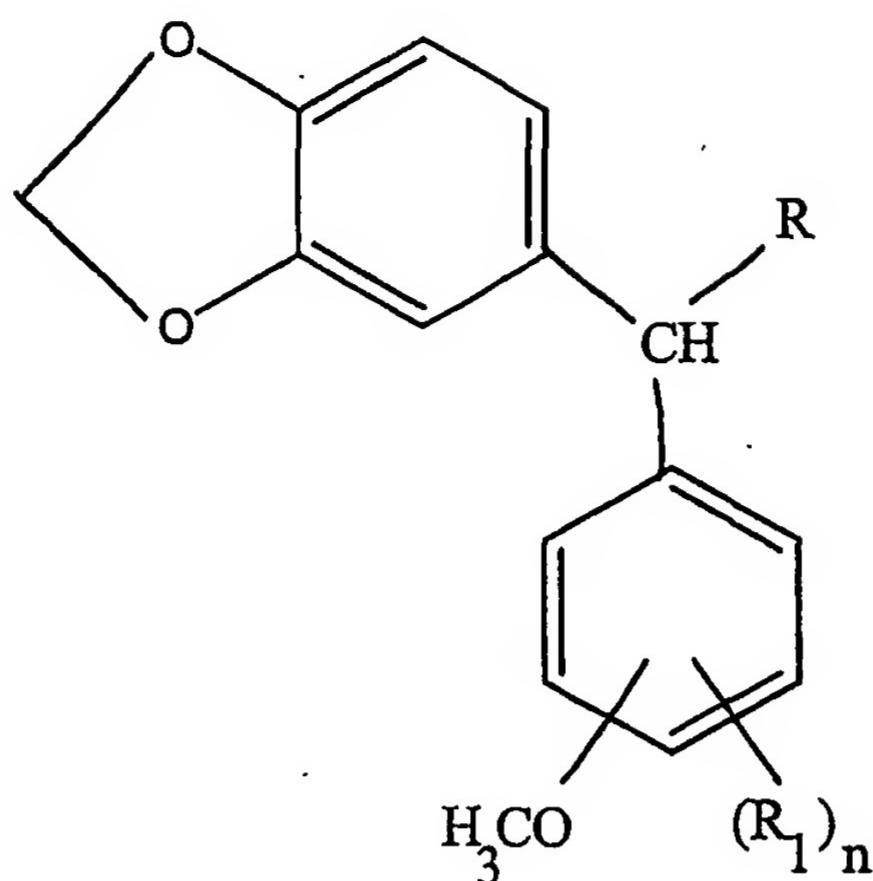
4. Use according to claim 1 of a compound of the formula III



5 wherein R₂, R₅, R₆, which can be the same or different, are H, OH, OOCCH₃, OOCH₂CH₃, OCH₃, or OC₂H₅, or R₅ and R₆ together is an ether or a lactone; and R₁ and n are as defined in claim 1, with the proviso that when R₂ is OH this substituent can not be in alpha-position.

10 5. Use according to claim 4 of a compound selected from group consisting of epipodophyllotoxin, deoxypodophyllotoxin, and acetylpodophyllotoxin.

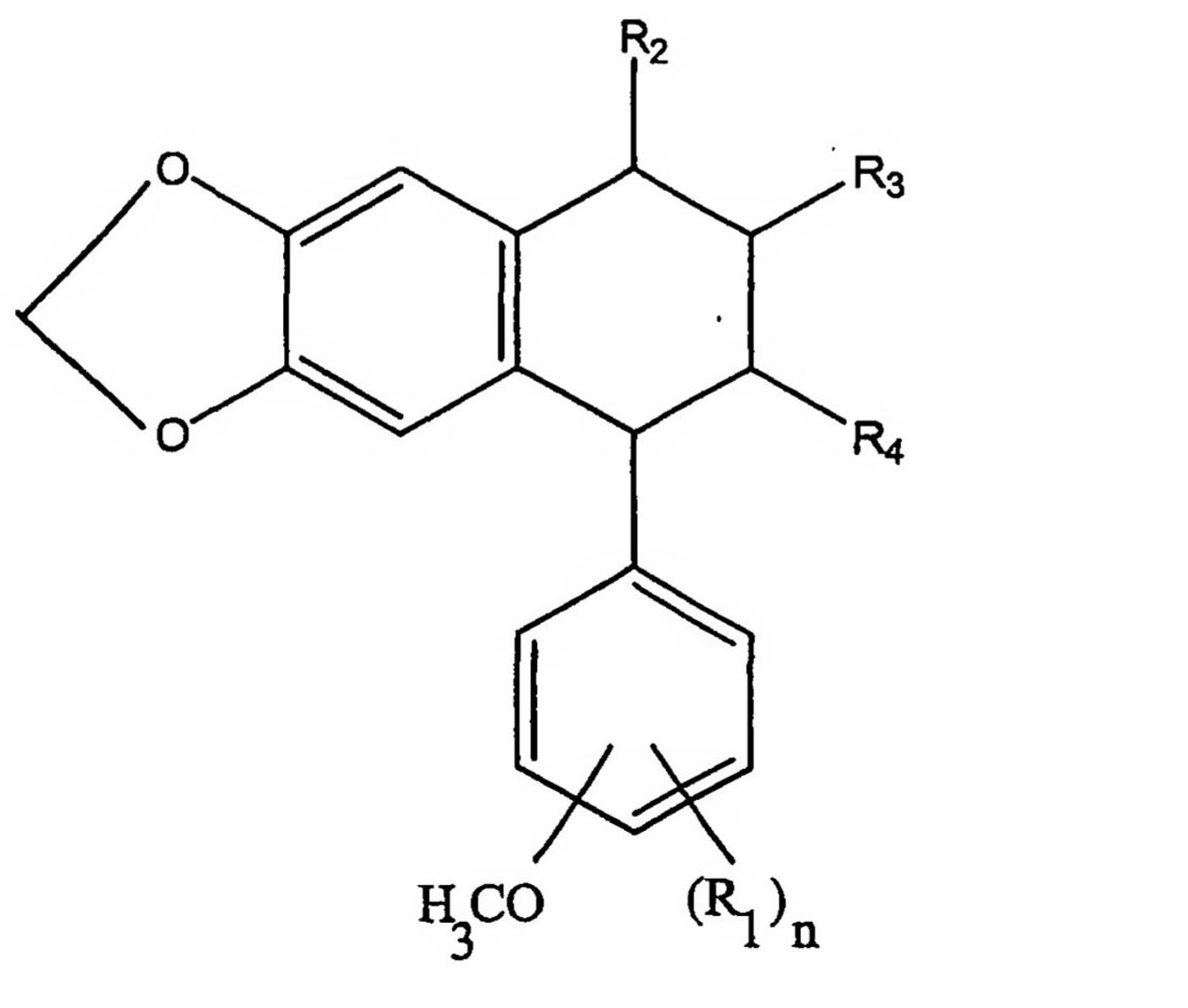
6. A compound of the formula I



5 wherein R is OH, OCH₃, OC₂H₅, CH₃, C₂H₅, or C₂H₄OH; and R₁ and n are as defined in claim 1, with the proviso that when R₁ is OCH₃ and n is 2, R is not OH.

7. A compound of the formula II

10



wherein R₂, R₃ and R₄, which can be the same or different, are H, OH, O, OCH₃, or R₂ and R₃ together is a methylenedioxy group, or R₃

and R₄ together is an acetonide, carbonate or methylenedioxy group; and R₁ and n are as defined in claim 1.

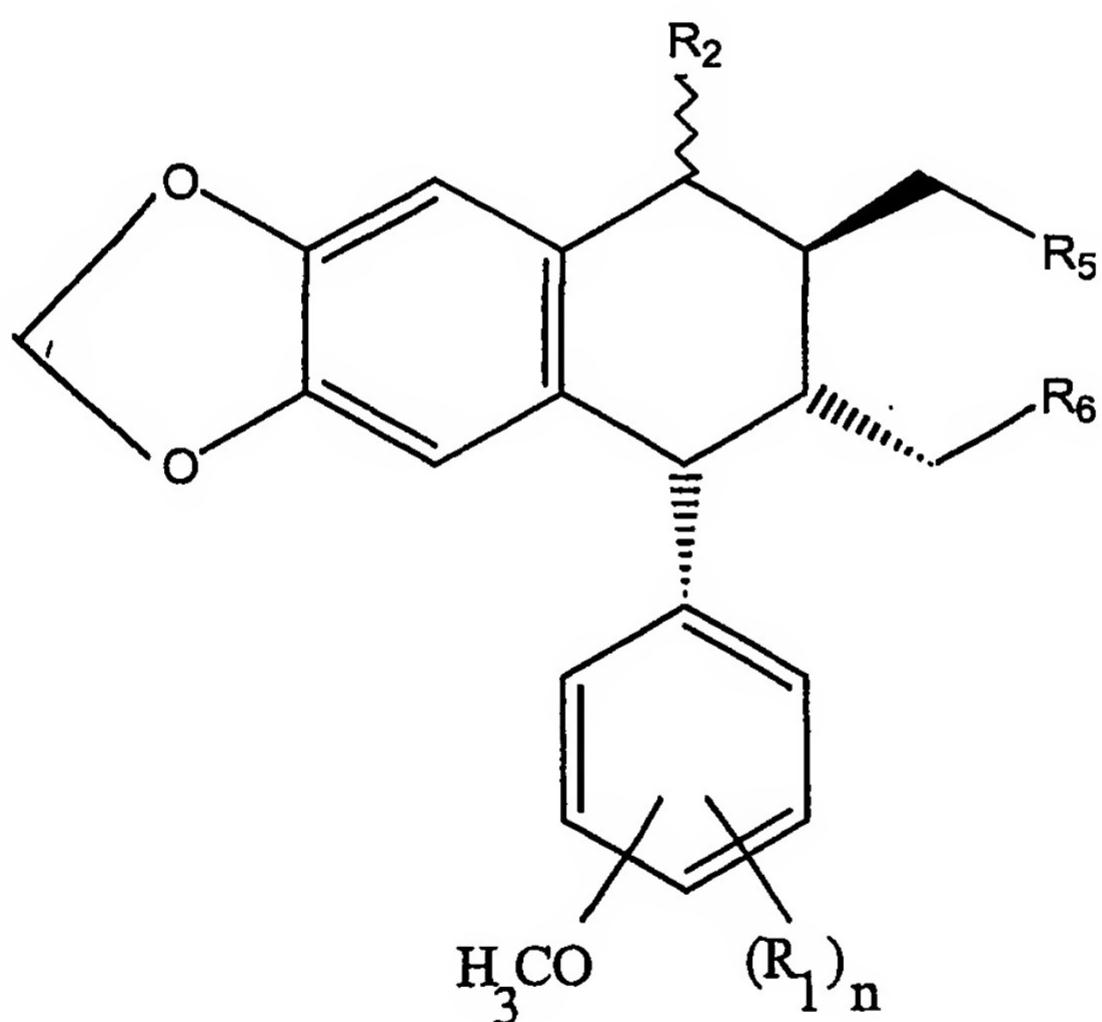
8. A compound of the formula I, II or III for use as a medicament.

5

9. Use of a compound of the formula I, II or III for treatment of IGF-1R dependent diseases, such as cancer, arteriosclerosis, psoriasis and acromegaly.

10 10. Use of a compound according to claim 6 or 7 for the preparation of a medicament for treatment of cancer.

11. Use of a compound of the formula III



15 III

wherein R₂, R₅, R₆, which can be the same or different, are H, OH, OOCCH₃, OOCH₂CH₃, OCH₃, or OC₂H₅, or R₅ and R₆ together is an ether or a lactone; and R₁ and n are as defined in claim 1, with the proviso that when R₂ is OH this substituent can not be in alpha-position,

20 for the preparation of a medicament for prophylaxis or treatment of cancer.

12. Use according to claim 11 of a compound selected from group consisting of epipodophyllotoxin, deoxypodophyllotoxin, and acetylpodophyllotoxin for prophylaxis or treatment of leukemia.

5 13. A pharmaceutical composition comprising a compound of the formula I, II or III in combination with a physiologically acceptable carrier.

10 14. Method of treatment of a cancer in a mammal, comprising the steps of administrating a pharmaceutical composition, containing a compound having the formula I, II or III in combination with a physiologically acceptable carrier, by constant infusion to a patient suffering from a tumor, monitoring the plasma level of the compound, and adjusting the rate of infusion to keep the plasma 15 level at a concentration of 0.05-5.0 μM , for a period of time being sufficient for the tumour to be retarded or to disappear.

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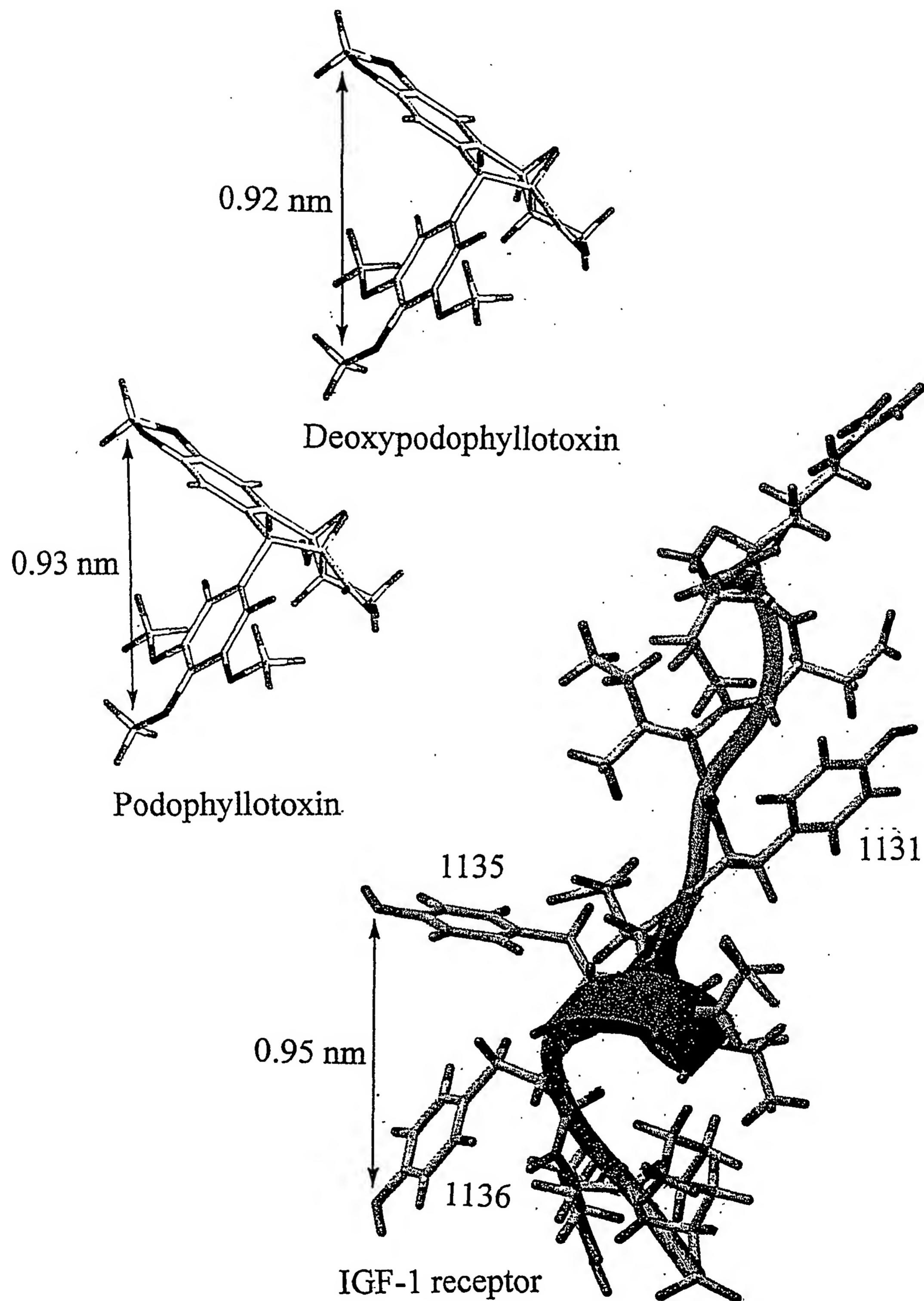


FIGURE 1

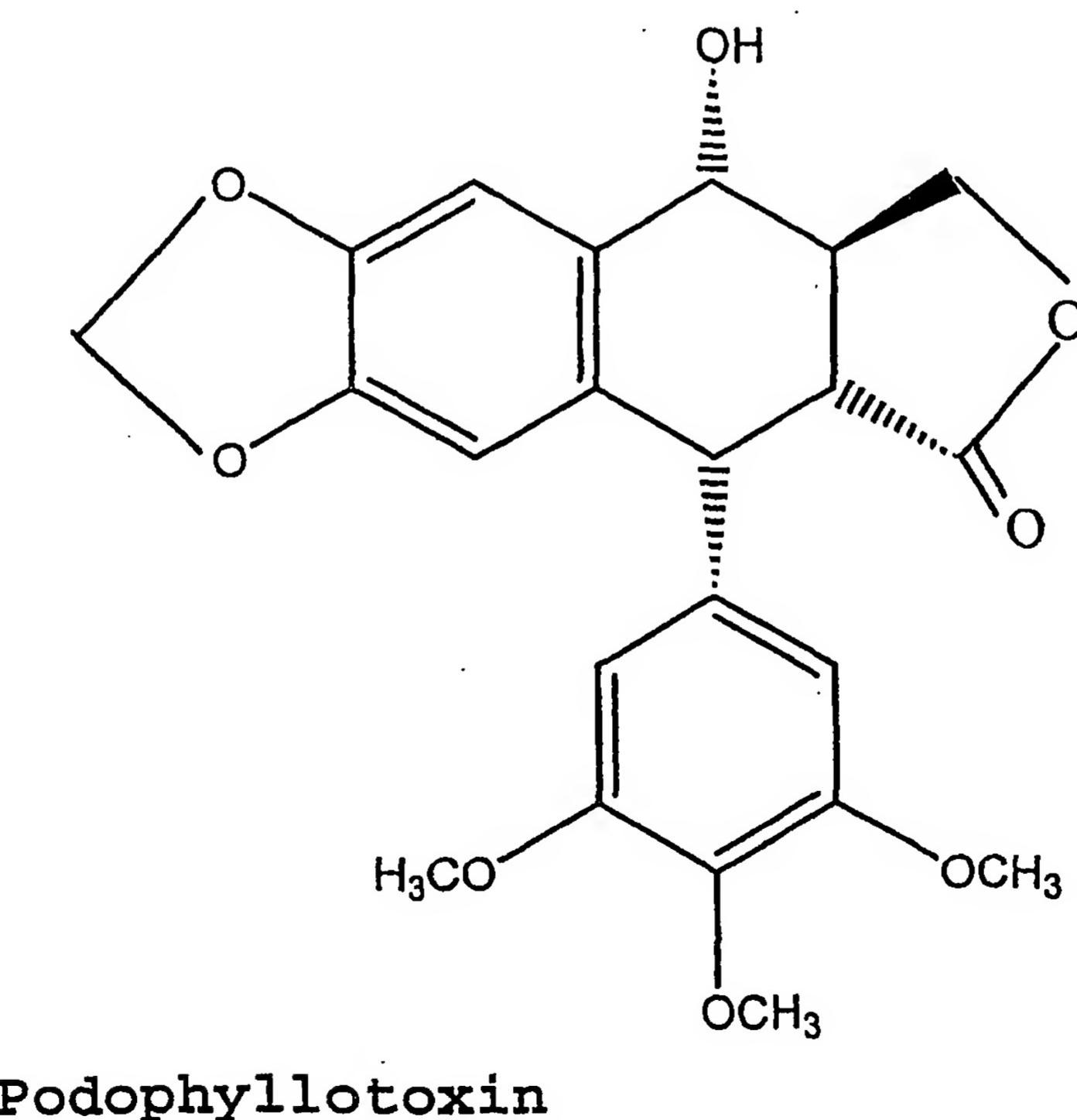
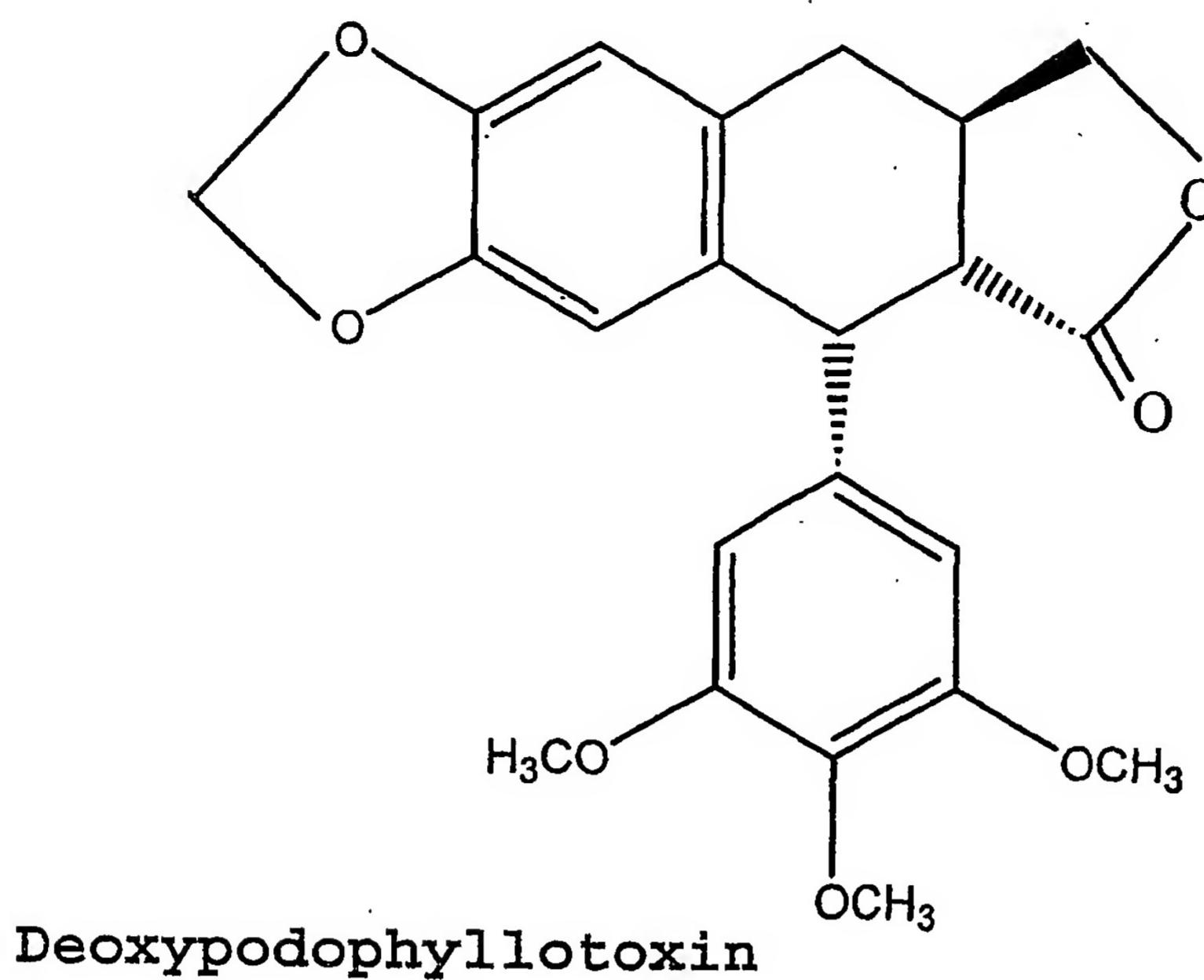
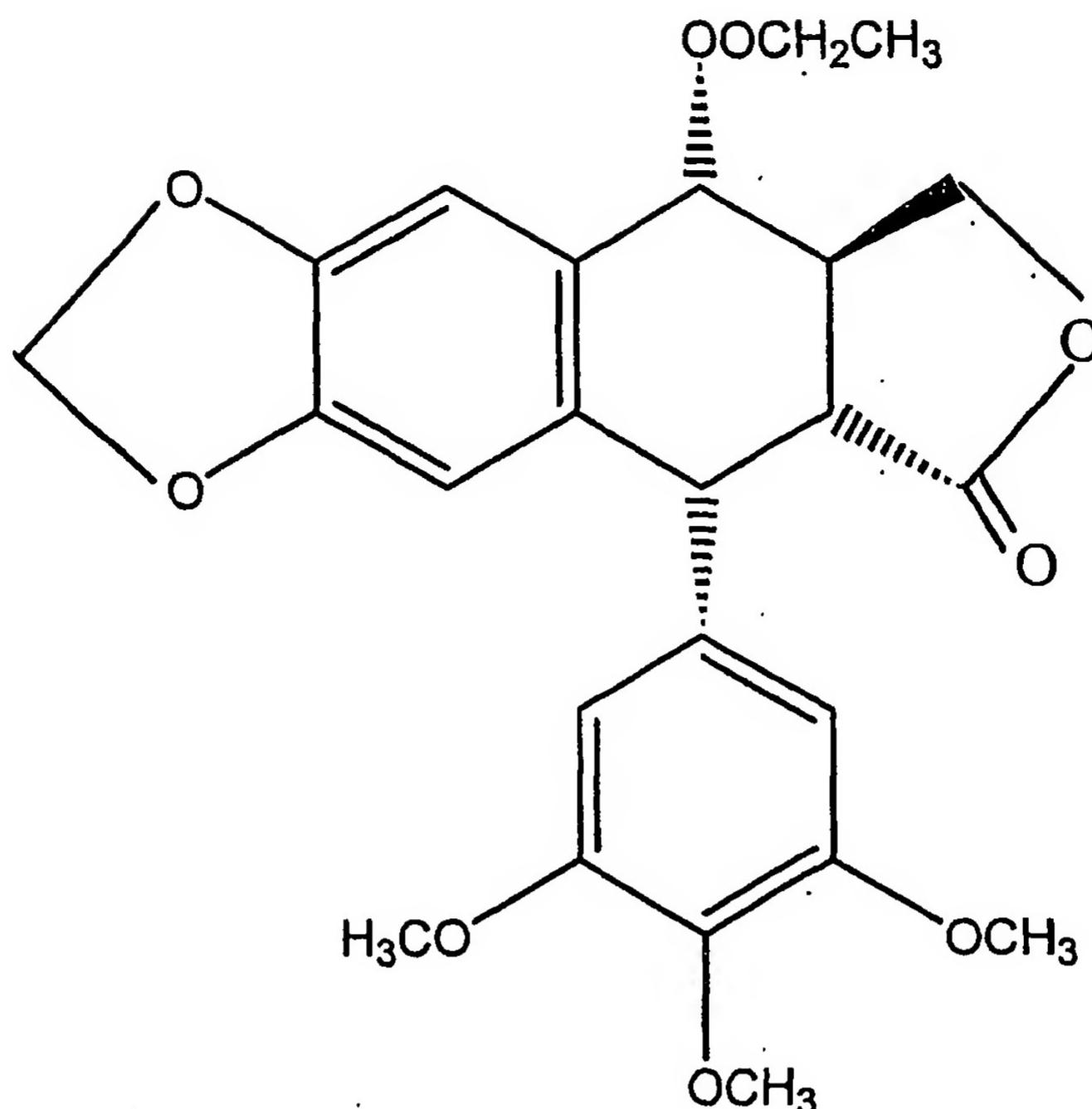
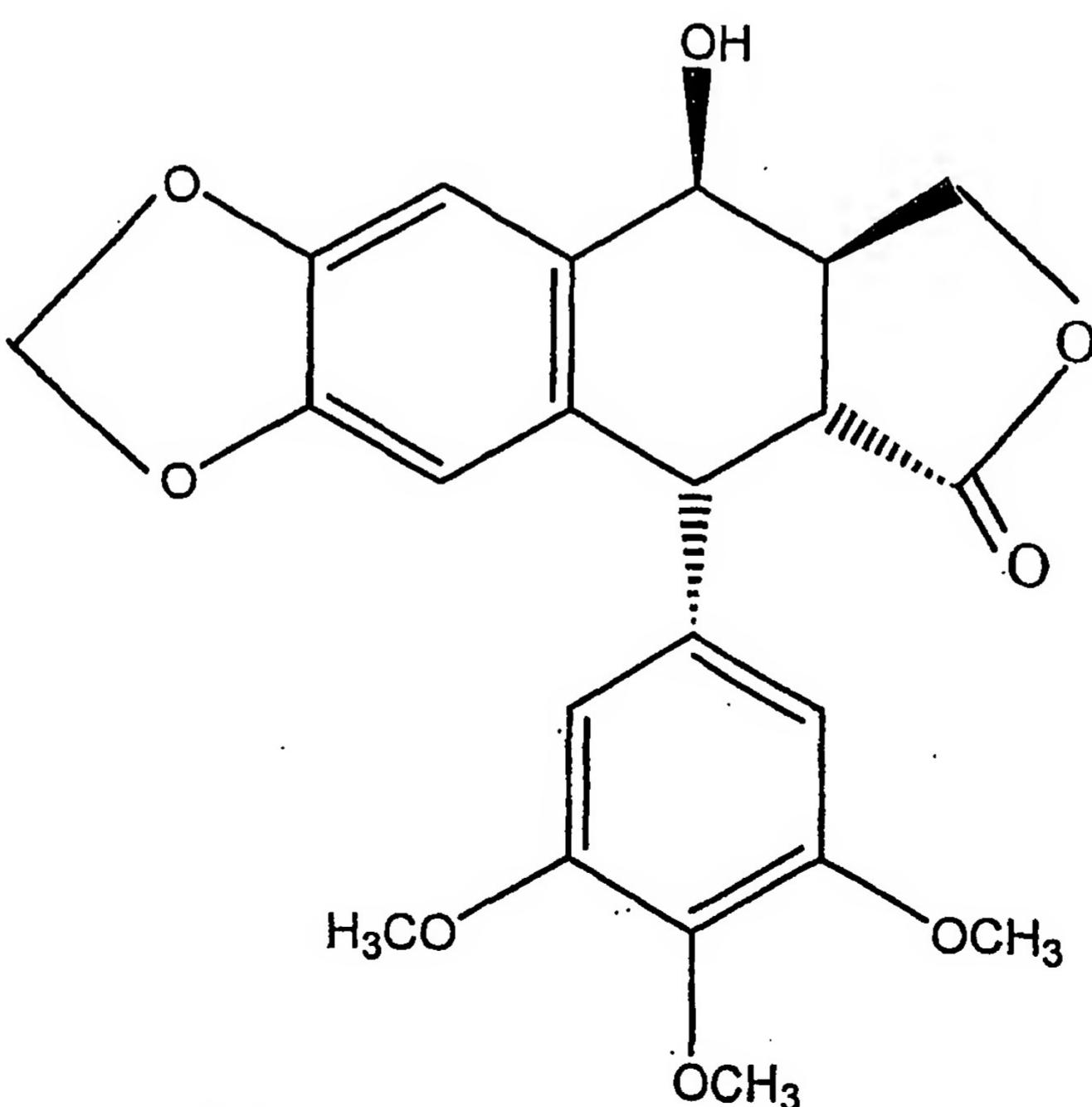


FIGURE 2A



Acetylpodophyllotoxin



Epipodophyllotoxin

FIGURE 2B

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01223

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C07D 493/04, C07D 317/48, A61K 31/36, A61K 31/34, A61P 5/06, A61P 9/10,
A61P 17/06, A61P 35/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C07D, A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-INTERNAL, WPI DATA, PAJ, CHEM. ABS DATA

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	WO 0240489 A1 (KIM, SONG-BAE), 23 May 2002 (23.05.02) --	1-14
X	Planta Medica, Volume 59, no. 3, 1999, A. San Feliciano et al: "Antineoplastic and Antiviral Activities of Some Cyclolignans", page 246 - page 249 --	1-14
X	Journal of Pharmaceutical Sciences, Volume 75, no. 11, November 1986, Ole Buchardt et al: "Thermal Chemistry of Podophyllotoxin in Ethanol and a Comparison of the Cytostatic Activity of the Thermolysis Products", page 1076 - 1080 --	1-14



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"E" earlier application or patent but published on or after the international filing date	"Y"	document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"&"	document member of the same patent family
"O" document referring to an oral disclosure, use, exhibition or other means		
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

18 Sept 2002

Date of mailing of the international search report

30 -09- 2002

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01223

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Arch. Pharm. (Weinheim), Volume 327, no. 3, 1994, Marina Gordaliza et al: "Antineoplastic and Antiviral Activities of Podophyllotoxin Related Lignans", page 175 - page 179	1-14
X	Journal of Natural Products, Volume 56, no. 10, October 1993, Miriam Novelo et al: "Cytotoxic constituents from hyptis verticillata", page 1728 - page 1736	1-14
X	Journal of Natural Products, Volume 58, no. 6, June 1995, J.M. Miguel del Corral: "Methyl ethers of podophyllotoxin-related cyclolignans", page 870 - page 877	1-14
X	Eur. J. Med. Chem., Volume 35, no. 7&8, 2000, Marina Gordaliza et al: "Synthesis and antineoplastic activity of cyclolignan aldehydes", page 691 - page 698	1-14
X	J. Med. Chem., Volume 39, no. 14, 1996, Marina Gordaliza et al: "Immunosuppressive Cyclolignans", page 2865 - page 2868	1-14
X	Tetrahedron, Volume 53, no. 46, 1997, Marina Gordaliza et al: "Preparation and Cytotoxicity of Podophyllotoxin Derivatives Lacking the Lactone Ring", page 15743 - page 15760	1-14
X	Bioorganic & Medicinal Chemistry Letters, Volume 5, no. 21, 1995, Marina Gordaliza et al: "Selective cytotoxic cyclolignans", page 2465 - page 2468	1-14

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01223

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0188248 A2 (TANABE SEIYAKU CO., LTD), 23 July 1986 (23.07.86), page 48 - page 51, page 4, line 11 - line 19, claims 8-23 --	1-14
X	WO 8604062 A1 (CONPHARM AB), 17 July 1986 (17.07.86), claims 1-4,6 --	1-14
X	SE 468213 B (CONPHARM AB), 23 November 1992 (23.11.92) --	1-14
X	PATENT ABSTRACTS OF JAPAN vol. 1997, no. 11, 28 November 1997 (1997-11-28) & JP 09194368 A (POLA CHEM IND INC), 29 July 1997 (1997-07-29) abstract --	1-14
X	US 2825730 A (EDWARD A. PRILL), 4 March 1958 (04.03.58), examples nos. 4, 5, 26 --	6
X	US 4342777 A (LEONARD JURD), 3 August 1982 (03.08.82), the claims --	6
X	STN International, File CAPLUS, CAPLUS accession no. 1985:125095, Document no. 102:125095, Batra, Janendra K. et al: "Structure-function studies with derivatives of 6-benzyl-1,3- benzodioxole, a new class of synthetic compounds which inhibit tubulin polymerization and mitosis"; & Mol. Pharmacol. (1985), 27(1), 94-102 --	1-14

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 02/01223

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	STN International, File CAPLUS, CAPLUS accession no. 1987:575915, Document no. 107:175915, Jurd, Leonard et al: "In vivo antitumor activity of 6-benzyl-1,3-benzodioxole derivatives against the P388, L1210, B16, and M5076 murine models"; J. Med. Chem. (1987), 30(10), 1752-6 --	1-14
A	WO 9928347 A1 (COMMONWEALTH SCIENTIFIC AND INDUSTRIAL RESEARCH ORGANISATION), 10 June 1999 (10.06.99) --	1-14
P,A	WO 0152826 A2 (SUPRATEK PHARMA INC.), 26 July 2001 (26.07.01) -- -----	1-14

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/SE 02/01223

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO	0240489	A1	23/05/02	NONE
EP	0188248	A2	23/07/86	
			SE 0188248 T3	
			AT 54441 T 15/07/90	
			AU 584153 B 18/05/89	
			AU 5175185 A 17/07/86	
			BG 45383 A 15/05/89	
			BG 45385 A 15/05/89	
			CN 1006464 B 17/01/90	
			CN 86100090 A 20/08/86	
			DD 270529 A 02/08/89	
			DE 3672501 D 00/00/00	
			ES 550578 A 16/05/87	
			ES 557052 A 16/12/87	
			ES 8705837 A 01/08/87	
			ES 8801239 A 01/03/88	
			FI 87557 B,C 15/10/92	
			FI 860089 A 11/07/86	
			HU 42428 A 28/07/87	
			HU 196737 B 30/01/89	
			IL 91117 A 10/03/91	
			JP 1839226 C 25/04/94	
			JP 2072136 A 12/03/90	
			JP 2072170 A 12/03/90	
			JP 5049668 B 26/07/93	
			JP 61267541 A 27/11/86	
			KR 8901884 B 29/05/89	
			NO 170760 B,C 24/08/92	
			NO 855355 A 11/07/86	
			PH 22105 A 01/06/88	
			PT 81814 A,B 01/02/86	
			US 4771072 A 13/09/88	
			US 4897418 A 30/01/90	
			US 5070103 A 03/12/91	

INTERNATIONAL SEARCH REPORT

International application No.
PCT/SE02/01223

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: **1–5, 9, 14**
because they relate to subject matter not required to be searched by this Authority, namely:
see next sheet
2. Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

Claims 1-5,9,14 relate to methods of treatment of the human or animal body by surgery or by therapy/ diagnostic methods practised on the human or animal body/Rule 39.1.(iv). Nevertheless, a search has been executed for these claims. The search has been based on the alleged effects of the compounds/compositions.

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